(54) Title: SUPPRESSORS OF CpG OLIGONUCLEOTIDES AND METHODS OF USE

(57) Abstract: The present disclosure relates to oligodeoxynucleotides that suppress an immune response. Methods are disclosed for preventing or treating an immune-mediated disorder, such as, but not limited to, an autoimmune disease, by administering a therapeutically effective amount of a suppressive oligodeoxynucleotide. Also disclosed are methods of suppressing an immune response in a subject by administering a therapeutically effective amount of a suppressive oligodeoxynucleotide.

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SUPPRESSORS OF CpG OLIGONUCLEOTIDES AND METHODS OF USE

PRIORITY CLAIM

This application claims the benefit of U.S. Provisional Patent Application

5   No. 60/324,484, filed September 24, 2001, and U.S. Provisional Patent Application

No. 60/400,826, filed August 1, 2002, which are both incorporated by reference

herein in their entirety.

FIELD OF THE DISCLOSURE

The present disclosure relates to oligodeoxynucleotides that suppress an

10   immune response, and to methods of using these oligonucleotides to treat disorders

associated with an immune response.

BACKGROUND

The immune system is composed of many interdependent cell types that

15   collectively protect the body from bacterial, parasitic, fungal, viral infections and

from the growth of tumor cells. Many of these cell types, such a B cells,

macrophages, an Natural Killer cells, have specialized functions. The cells of the

immune system can engulf bacteria, kill parasites or tumor cells, or kill viral-

infected cells. Often, these cells depend on the T helper subset for activation signals

20   in the form of secretions formally known as cytokines, lymphokines, or more

specifically interleukins.

Cells of the immune system recognize and are activated by conserved

pathogen associated molecular patterns (PAMPs) in infectious agents. The

unmethylated CpG dimers embedded in bacterial DNA, as well as certain synthetic

25   oligodeoxynucleotides (ODNs) containing unmethylated CpG sequences (termed a

CpG motif) that emulated them, are more frequent in the genomes of bacteria and

viruses than vertebrates. Recent studies suggest that immune recognition of these

motifs may contribute to the host's innate immune response (Klinman *et al.*, *Proc.*

Natl. Acad. Sci. USA 93: 2879, 1996; Yi *et al.*, *J. Immun.* 157: 5394, 1996; Liang *et*

30   *al.*, *J. Clin. Invest.* 98:1119, 1996; Krieg *et al.*, *Nature* 374: 546, 1995).

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In mice, CpG DNA induces proliferation in almost all (>95%) of B cells and increases immunoglobulin (Ig) secretion. This B-cell activation by CpG DNA is T-cell independent and antigen non-specific. In addition to its direct effects on B cells, CpG DNA has also been shown to activate cells of the immune system (see, for

5 example, International Patent Applications WO 95/26204, WO 96/02555, WO 98/11211, WO 98/18810, WO 98/37919, WO 98/40100, WO 98/52581, PCT/US98/047703, and PCT/US99/07335; U.S. Patent No. 5,663,153).

However, in many situations, there is a need to suppress an immune response. For example, in an autoimmune disease, a foreign antigen mimics one or

10 more self-proteins, and the immune system produces a response in which a tissue is consequently injured. Similarly, when a subject is the recipient of a transplanted tissue (*e.g.* a heart, lung, pancreas, or kidney recipient), the body can produce an immune response against the donor tissue. In this situation, there is a clear need to suppress the immune response, in order to avoid rejection of the graft. Additionally,

15 there is a need to suppress the immune response in order to prevent or treat allergic disorders such as asthma.

In view of the above, there exists a need for agents that suppress immune responses. Specifically, there is a need for agents that can be used to suppress the inflammation, and that can be used to suppress an the immune response associated

20 with autoimmune diseases, allergies, and transplant rejection.

BRIEF SUMMARY OF SPECIFIC EMBODIMENTS

Oligodeoxynucleotides are disclosed herein that can be used to suppress immune activation. These suppressive oligodeoxynucleotides are of use in

25 preventing and/or treating a variety of diseases and disorders that include, but are not limited to, autoimmune diseases. Specific, non-limiting examples of autoimmune disorders are inflammatory arthritis, Hashimoto's thyroiditis, pernicious anemia, Addison's disease, type I diabetes, systemic lupus erythematosus, dermatomyositis, Sjogren's syndrome, dermatomyositis, lupus

30 erythematosus, multiple sclerosis, myasthenia gravis, Reiter's syndrome, and Grave.s disease, among others. The suppressive oligodeoxynucleotides can be

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administered locally, such as, but not limited to, administration by intra-articular injection or inhalation, or can be administered systemically.

A substantially pure or isolated oligodeoxynucleotide (ODN) is disclosed herein that is at least about 8 nucleotides in length, forms a G tetrad, has a CD value

5   of greater than 2.9, has at least two guanosines, and suppresses an immune response. Optionally, the suppressive ODN has multiple guanosine-rich sequences, and in some examples, the ODN has one or more TTAGGG motifs. Furthermore, in particular embodiments, the ODN is modified to prevent degradation or is part of an oligodeoxynucleotide delivery complex that includes a targeting moiety. In one

10   specific, non-limiting example the suppressive ODN suppresses CpG-DNA-induced immune activation.

Also disclosed herein is a pharmacological composition that includes the suppressive ODN and a pharmacologically acceptable carrier.

In another embodiment, a method is disclosed for treating or preventing an

15   autoimmune disease in a subject. The method includes administering a therapeutically effective amount of the suppressive ODN to a subject having or at risk of developing an autoimmune disease, thereby treating or preventing the autoimmune disease. In some embodiments, the ODN is administered orally, intravenously, intra-muscularly, sub-cutaneously, or intra-articularly.

20   In another embodiment, a method is disclosed for treating or preventing an autoimmune disease in a subject that includes contacting immune cells with the suppressive ODN *in vitro*. The immune cells are and transferred to a subject having or at risk of developing an autoimmune disease.

Other embodiments are methods of suppressing an immune response in a

25   subject including administering a therapeutically effective amount of a suppressive ODN to a subject in which it is desirable to suppress an immune response, thereby suppressing the immune response.

Also described herein is a kit for treating or preventing an autoimmune disease in a subject that includes the suppressive ODN and instructions for

30   administering the ODN to a subject.

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BRIEF DESCRIPTION OF THE FIGURES

Fig. 1: is a set of diagrams of the structure of a G-tetrad. Fig. 1A is a
diagram of the structure of an individual G-tetrad that shows the Hoogsteen base
5   pairing. M^+ represents a monovalent cation such as K^+ or Na^+ and dR is the sugar-
phosphate backbone. Fig. 1B is a schematic representation showing the possible
folded intramolecular quadruplex structure. Fig. 1C is a schematic showing the GG-
base pair formed by means of Hoogsteen hydrogen bonds. Fig. 1D is a schematic of
an intramolecular hairpin.
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Fig. 2: is a set of graphs showing the factors contributing to the suppression
of CpG-induced immune activation. Fig. 2A is a graph showing that mammalian
DNA suppresses CpG DNA-induced immune activation. Fig. 2B is a graph
showing that the telomeric TTAGGG repetitive motif is suppressive. Fig. 2C is a
15   graph showing that the suppressive motif is active in *trans* and *cis* conformations.
Fig. 2D is a graph showing that suppressive ODNs selectively block CpG DNA-
induced immune activation. Fig. 2E is a graph showing that Poly Gs are critical for
suppression. Fig. 2F is a graph showing that G-tetrad forming non-telomere
sequences are also suppressive.
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Fig. 3: is a pair of graphs showing that G-tetrad-forming suppressive ODNs
selectively suppress CpG-induced immune activation. Fig. 3A is a graph showing
that oligonucleotides (ODNs) containing suppressive motifs inhibit CpG-induced
immune activation in a dose dependent fashion. Fig. 3B is a graph showing G-tetrad
25   formation and suppressive activity of phosphorothioate and 7-DG modified ODNs.

Fig. 4: is a graph showing the effect of suppressive ODN on CpG DNA and
Con A induced IFNγ production. BALB/c spleen cells were stimulated with 1 μM
CpG ODN (ODN1555, ODN1466), 50 μg/ml of bacterial DNA, or 5 μg/ml Con A.
30   The response of these cultures was compared to cells co-stimulated with 1 μM of
control ODN1612, suppressive ODN1502 or suppressive ODNH154. The number
of IFNγ secreting cells was determined by ELIspot after 18 h. Data represent the

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average + SD of triplicate cultures. The experiment was repeated three times with similar results.

Fig. 5: is a graph showing the concentration effects of suppressive ODN.
5   BALB/c spleen cells were stimulated with 1 μM CpG ODN1555 or ODN1466 plus
increasing amounts of suppressive ODN1502 or ODNH154. Cytokine levels in
culture supernatants were measured by ELISA after 24 h. Results represent the mean
+ SD of 4 different experiments.

10   **Fig. 6:** is a graph showing the kinetics of suppressive ODN. BALB/c spleen
cells were stimulated with 1 μM CpG ODN1555. At various times, 1 μM
suppressive ODN1502 was added. Cytokine levels in culture supernatants were
measured by ELISA after 24 h. Results represent the mean of two independent
experiments.
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Fig. 7: is a graph showing the effect of removing CpG ODN from cultured
cells. 1 μM of CpG ODN1555 was added to BALB/c spleen cells at T = 0. The
cells were washed free of this ODN after various incubation periods. IFNγ and IL-
12 levels in culture supernatants were measured by ELISA after 24 h. Results
20   represent the average + SD of duplicate cultures. Similar results were obtained in
studies of CpG ODN1466.

Fig. 8: is a graph showing that suppressive ODN do not block the binding or
uptake of CpG ODN. BALB/c spleen cells were incubated with 1 μM of CpG
25   ODN1555 (black bar) plus 1 μM of suppressive ODN1502 (grey bar) or control
ODN1612 (white bar). The percent of cells that bound or internalized the CpG ODN
was determined by FACS. Similar results were obtained using CpG ODN1466,
suppressive ODNH154 and control ODN1471.

30   **Fig. 9:** is a graph showing that suppressive ODN permit higher levels of
transgene expression.

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Fig. 10: is a graph showing that suppressive ODN blocks the activity of CpG motifs *in vivo*.

Fig. 11: is a set of graphs showing the effect of CpG ODN and suppressive
5   ODN injection into the knee. Fig. 11A is a graph of joint swelling in mice treated with CpG ODN (solid circle), CpG plus control ODN (solid square), CpG plus suppressive ODN (open diamond), control ODN (open triangle), suppressive ODN (inverted open triangle), or PBS (open square, N = 8-11 mice/group). Fig. 11B is a graph showing histologic changes in the injected knees. Knees were evaluated 4
10 days after treatment by a blinded investigator. Scale: 0; no inflammation, 1; sparse, localized perivascular infiltrate, 2; moderate infiltrate, 3; moderate – dense infiltrate with synovial hyperplasia. Fig. 11C is a graph showing that suppressive ODNs suppress joint swelling. At T=0, both knees were injected with 25 g of CpG ODN. The R knee was then injected with PBS and the L knee with 25 g of suppressive
15 ODN either 0 (solid circles, N=3), 24 (solid squares, N=6) or 48 (solid triangles, N=6) h later. Results show the difference in swelling between the two knees. Statistical significance was assessed by repeated-measures ANOVA using the Proc Mixed procedure (A, C) and Wilcoxon Rank sum test (B). * $p<0.05$.

20 Fig. 12: is a digital image of a photomicrograph showing the effects of CpG and ODN injection on knee histology in mice. BALB/c knees were injected with PBS (Fig. 12A, B), CpG ODN (Fig. 12C, D), CpG plus control ODN (Fig. 12E, F) or CpG plus suppressive ODN (25 g of each ODN, Fig. 12G, H). Typical histology 4 days after injection of 25 μg of each ODN is shown. Note the cellular infiltrates,
25 perivascular accumulation of mononuclear cells, and hyperplasia of the synovial lining in the knees of mice injected with CpG ODN. Panels A, C, E and G show 100x magnification; panels B, D, F, H show 400x magnification.

Fig. 13: is a set of graphs that demonstrate that administration of
30 suppressive ODN decreases TNFα upregulation following CpG ODN injection. Fig. 13A is a graph of TNFα levels following treatment with ODN. BALB/c spleen cells were stimulated *in vitro* for 72 h with 1 μM of various ODN. Fig. 13B is a graph of

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percent TNFα production versus suppressive ODN concentration. RAW 264.7 cells (106/well) were stimulated with 1 uM CpG plus increasing amounts of suppressive ODN. The concentration of TNF in culture supernatants after 24 h was measured by ELISA. Data represent the mean + SEM of 5 independently studied animals/group.

5   Statistical significance was assessed by Wilcoxon Rank sum test. Fig. 13C is a digital image of an agarose gel. Joints injected with 25 g of ODN were processed into RNA 3 days later. Representative examples of local TNF and β-actin mRNA levels are shown. Fig. 13D is a graph of relative TNFα mRNA expression following treatment with CpG ODN or CpG ODN and suppressive ODN. Relative intensity of

10   TNFα vs β-actin mRNA (N=3). * p<0.05.

       Fig. 14: is a graph showing that the local administration of suppressive ODNs reduces the pro-inflammatory effect of CpG ODN administration. Optimal suppression of CpG ODN-mediated joint inflammation is attained when the

15   suppressive ODN are administered 3 days before the inflammatory challenge. Shown is the mean and SEM of 10-11 mice/group pre-treated locally (intra-articularly) with suppressive ODN 3 days prior to CpG ODN challenge (open circles), suppressive ODN administered 1 day prior to CpG ODN challenge (open, inverted triangles), suppressive ODN administered at the time of the CpG ODN

20   challenge (open triangles). The joint swelling of mouse knees injected with control ODN (black diamonds) or PBS (black circles) 3 days before the CpG serve as controls.

       Fig. 15: is a graph showing that naive mice (and mice pre-treated with

25   control ODN or PBS) developed severe arthritis following local CpG ODN challenge. In contrast, systemic administration of suppressive ODN three days prior to local CpG DNA challenge reduced joint swelling and inflammation by 80 - 85% (p<0.029). BALB/c mice were treated IP with 300 µg of suppressive ODN 0-3 days prior to the intra-articular injection of 25 µg CpG DNA. Consistent with previous

30   studies, naive mice (and mice pre-treated with control ODN or PBS) developed severe arthritis following local CpG ODN challenge.

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Fig. 16: is a pair of graphs showing that systemic administration of suppressive ODN three days prior to local CpG DNA challenge reduced joint swelling and inflammation by 80-85% (p<0.029).

5   Fig. 17: is a pair of graphs showing that systemically administered suppressive ODN elicit a population of regulatory cells that inhibit CpG-induced arthritis. As expected, spleen cells from untreated donors had no effect on the development of CpG-induced arthritis. By comparison, the transfer of 20 x 10^6 spleen cells from suppressive ODN treated donors significantly reduced joint

10   swelling and inflammation in the recipients.

Fig. 18: is a graph showing that CD11c+ cells are responsible for the resistance to CpG-induced arthritis. Magnetic beads were used to deplete or enrich specific cell subpopulations from donor spleens. Depletion of CD19+ B cells, T

15   cells, or NK cells had no effect on CpG-induced arthritis. However, removal of CD11c+ dendritic cells resulted in a complete abrogation of the suppressive activity of the transferred spleen cell population. Similarly, the transfer of only 5 x 105 CD11c+ enriched cells from suppressive ODN treated mice to normal recipients conferred resistance to CpG- induced arthritis.

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Fig. 19. is a graph showing that neutralizing ODNs inhibit HSV DNA-induced angiogenesis. Pellets containing 1 μg of HSV DNA alone or in combination with 1 μg of control or neutralizing ODN were implanted into corneal micropockets. The figure shows the average degree of angiogenesis 4 days after

25   pellet implantation (four to five mice per group). *, $P < 0.01$.

BRIEF DESCRIPTION OF THE SEQUENCE LISTING

SEQ ID NOs 1-25 are suppressive ODN sequences.

SEQ ID NOs 28, 29, and 31 are control ODN sequences.

30   **SEQ ID NOs 26, 27, and 30** are immunostimulatory CpG sequences.

DETAILED DESCRIPTION OF SEVERAL EMBODIMENTS

I. *Abbreviations*

A: adenine

5   Ab: antibody

C: cytosine

CD: circular dichroism

CpG ODN: an oligodeoxynucleotide including a CpG motif.

DC: dendritic cell

10   FCS: fetal calf serum

G: guanine

GI: gastrointestinal

GU: genitourinary

h: hour

15   IFN-α: interferon alpha

IFN-γ: interferon gamma

IL-10: interleukin 10

mm: millimeter

mRNA: messenger ribonucleic acid.

20   ODN: oligodeoxynucleotide

Pu: purine

Py: pyrimidine

s.c.: subcutaneous

T: thymine

25   μg: microgram

II. *Terms*

Unless otherwise noted, technical terms are used according to conventional
usage. Definitions of common terms in molecular biology may be found in
30 Benjamin Lewin, *Genes V*, published by Oxford University Press, 1994 (ISBN 0-19-
854287-9); Kendrew *et al.* (eds.), *The Encyclopedia of Molecular Biology*, published

by Blackwell Science Ltd., 1994 (ISBN 0-632-02182-9); and Robert A. Meyers

(ed.), *Molecular Biology and Biotechnology: a Comprehensive Desk Reference*,

published by VCH Publishers, Inc., 1995 (ISBN 1-56081-569-8).

5   Unless otherwise explained, all technical and scientific terms used herein

have the same meaning as commonly understood by one of ordinary skill in the art

to which this disclosure belongs. The singular terms "a," "an," and "the" include

plural referents unless context clearly indicates otherwise. Similarly, the word "or"

is intended to include "and" unless the context clearly indicates otherwise. It is

further to be understood that all base sizes or amino acid sizes, and all molecular

10   weight or molecular mass values, given for nucleic acids or polypeptides are

approximate, and are provided for description. Although methods and materials

similar or equivalent to those described herein can be used in the practice or testing

of the present disclosure, suitable methods and materials are described below. In

case of conflict, the present specification, including explanations of terms, will

15   control. In addition, the materials, methods, and examples are illustrative only and

not intended to be limiting.

In order to facilitate review of the various embodiments of the disclosure, the

following explanations of specific terms are provided:

20   **Allergy:** An example of an immune-mediated disorder. An allergy is a

collection of symptoms caused by an exaggerated immune response or reaction to

substances that do not trigger an immune response in most people. The term

"allergy" has become synonymous with Type I hypersensitivity (IgE-mediated

allergy). Four different types of hypersensitivity were described by Coomb and Gell

25   (Types I, II, III and IV), as a pedagogical way to increase the understanding of

different immune reactions which could be provoked by many antigens. In practice

these types do not necessarily occur in isolation from each other.

Allergic diseases generally begin in childhood, although they can arise at any

age. Development of allergic disease is associated with an allergic constitution due

30   to heredity and to environmental and health factors. An allergic response involves

an increased production of allergen-specific IgE antibodies, which may lead to

clinical symptoms such as rhinitis, asthma, eczema, colic pains or diarrhea. A state

of hyperreactivity often accompanies an allergic reaction. If this hyperreactivity occurs in the respiratory tract, everyday stimuli like dust, tobacco smoke, cold air and perfumes may lead to allergy-like symptoms.

Animal: Living multi-cellular vertebrate organisms, a category that includes,
5   for example, mammals and birds. The term mammal includes both human and non-human mammals. Similarly, the term "subject" includes both human and veterinary subjects.

Angiogenesis: A process leading to the generation of new blood vessels through sprouting from already existing blood vessels. The process involves the
10   migration and proliferation of endothelial cells from preexisting vessels. Angiogenesis occurs both during pre-natal development, post-natal development, and in the adult. In the adult angiogenesis occurs during the normal cycle of the female reproductive system, wound healing, and during pathological processes such as cancer (for review see Battegay, *J. Molec. Med.* 73(7): 333-346, 1995; Beck and
15   D'Amore, *FASEB J.* 11(5): 365, 1997)

Angiogenic Factor: A molecule that promotes angiogenesis. A plethora of experiments have suggested that tissues secrete factors which promote angiogenesis under conditions of poor blood supply during normal and pathological angiogenesis processes. Angiogenic molecules are generated by tumor, inflammatory, and
20   connective tissue cells in response to hypoxia and other as-yet ill-defined stimuli. The first indication of the existence of such diffusible substances was gleaned from filtration experiments demonstrating that tumor cells separated from underlying tissues by filters that do not allow passage of cells are nevertheless capable of supporting vessel growth in these tissues. The formation of blood vessels is initiated
25   and maintained by a variety of factors secreted either by the tumor cells themselves or by accessory cells. Many different growth factors and cytokines have been shown to exert chemotactic, mitogenic, modulatory or inhibitory activities on endothelial cells, smooth muscle cell and fibroblasts and can, therefore, be expected to participate in an angiogenic process in one way or another. For example, factors
30   modulating growth, chemotactic behavior and/or functional activities of vascular endothelial cells include αFGF, βFGF, angigiogenein, angiotropin, epithelial growth factor, IL-8, and vascular endothelial growth factor (VEGF), amongst others.

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As many angiogenic factors are mitogenic and chemotactic for endothelial cells their biological activities can be determined *in vitro* by measuring the induced migration of endothelial cells or the effect of these factor on endothelial cell proliferation. Alternatively, a bioassay may be utilized for direct determination of
5   angiogenic activities and permit repeated, long-term quantitation of angiogenesis as well as physiological characterization of angiogenic vessels. Many such assays are known in the art.

One assay employs the use of a non-vascularized mouse eye (*e.g.* Kenyon *et al., Invest Opthalmol. Vis. Sci.* 37:1625, 1996; also see Examples section) or the
10   rabbit eye (*e.g.*, see Gaudric *et al,. Ophthal.* Res. 24: 181, 1992), and is termed a cornea pocket assay. This assay has the advantage that new blood vessels are easily detected and essentially must be newly formed blood vessels in the normally avascular cornea. Another assay involves the use of chicken chorioallantoic membrane (the CAM assay; see Wilting *et al., Anat. Embryol.* 183: 259, 1991).
15   Other assays in the rat, such as the rat aortic ring model, provide reproducible assays that are often utilized to identify angiogenic agonists and antagonists (*e.g.* see Lichtenberg *et al., Pharmacol Toxicol.* 84: 34, 1999).

Antigen: A compound, composition, or substance that can stimulate the production of antibodies or a T-cell response in an animal, including compositions
20   that are injected or absorbed into an animal. An antigen reacts with the products of specific humoral or cellular immunity, including those induced by heterologous immunogens. The term "antigen" includes all related antigenic epitopes.

Antisense, Sense, and Antigene: Double-stranded DNA (dsDNA) has two strands, a 5' -> 3' strand, referred to as the plus strand, and a 3' -> 5' strand (the
25   reverse compliment), referred to as the minus strand. Because RNA polymerase adds nucleic acids in a 5' -> 3' direction, the minus strand of the DNA serves as the template for the RNA during transcription. Thus, the RNA formed will have a sequence complementary to the minus strand and identical to the plus strand (except that U is substituted for T).
30   Antisense molecules are molecules that are specifically hybridizable or specifically complementary to either RNA or the plus strand of DNA. Sense molecules are molecules that are specifically hybridizable or specifically complementary to the

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minus strand of DNA. Antigene molecules are either antisense or sense molecules directed to a dsDNA target. In one embodiment, an antisense molecule specifically hybridizes to a target mRNA and inhibits transcription of the target mRNA.

Arthritis: Arthritis is an inflammatory disease that affects the synovial

5   membranes of one or more joints in the body. It is the most common type of joint disease, and it is characterized by the inflammation of the joint. The disease is usually oligoarticular (affects few joints), but may be generalized. The joints commonly involved include the hips, knees, lower lumbar and cervical vertebrae, proximal and distal interphangeal joints of the fingers, first carpometacarpal joints,

10   and first tarsometatarsal joints of the feet.

One type of arthritis is reactive arthritis, which is an acute nonpurulent arthritis secondary to a urinary tract or gastrointestinal infection with a variety of microorganisms, including *Chlamydia trachomatis, Yersinia, Salmonella, Shigella,* and *Campylobacter*. Microbial components are found in the affected joints. The

15   arthritis appears abruptly and tends to involve the knees and ankles, but sometimes involves the wrists, fingers, and/or toes. Untreated, the arthritis lasts for about a year, then generally abates and only rarely is accompanied by ankylosing spondylitis. Despite evidence of disease being triggered by bacterial infection, viable bacteria are rarely present in affected joints and antibiotic treatment seldom

20   provides relief.

Another type of arthritis is rheumatoid arthritis. Rheumatoid arthritis is a chronic, systemic, inflammatory disease that affects the synovial membranes of multiple joints in the body. Because the disease is systemic, there are many extra-articular features of the disease as well. For example, neuropathy, scleritis,

25   lymphadenopathy, pericarditis, splenomegaly, arteritis, and rheumatoid nodules are frequent components of the disease. In most cases of rheumatoid arthritis, the subject has remissions and exacerbations of the symptoms. Rheumatoid arthritis considered an autoimmune disease that is acquired and in which genetic factors appear to play a role.

30   **Autoimmune disorder:** A disorder in which the immune system produces an immune response (*e.g.,* a B cell or a T cell response) against an endogenous antigen, with consequent injury to tissues. For example, rheumatoid arthritis is an

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autoimmune disorder, as are Hashimoto's thyroiditis, pernicious anemia, Addison's disease, type I diabetes, systemic lupus erythematosus, dermatomyositis, Sjogren's syndrome, dermatomyositis, lupus erythematosus, multiple sclerosis, myasthenia gravis, Reiter's syndrome, and Grave.s disease, among others.

5   **CD value:** The formation of G-tetrads yields a complex with different physical properties than the individual oligonucleotides. Spectroscopically, this is manifested by an increase in circular dicroism (**CD**), and an increase in peak absorbance to the 260-280 nm wavelength owing to the formation of secondary structures. Thus, a convenient method for identifying oligonucleotides that form G-

10   tetrads is to study their **CD values.** An increase in peak ellipticity values to greater than 2.0 is typical of a G-tetrad forming oligonucleotide. The higher the ellipticity value, the greater the tetrad-forming capacity of the oligonucleotide.

       CpG or CpG motif: A nucleic acid having a cytosine followed by a guanine linked by a phosphate bond in which the pyrimidine ring of the cytosine is

15   unmethylated. The term "methylated CpG" refers to the methylation of the cytosine on the pyrimidine ring, usually occurring the 5-position of the pyrimidine ring. A CpG motif is a pattern of bases that include an unmethylated central CpG surrounded by at least one base flanking (on the 3' and the 5' side of) the central CpG. Without being bound by theory, the bases flanking the CpG confer part of the

20   activity to the CpG oligodeoxynucleotide. A CpG oligonucleotide is an oligonucleotide that is at least about ten nucleotides in length and includes an unmethylated CpG. CpG oligonucleotides include both D and K type oligodeoxynucleotides (see below). CpG oligodeoxynucleotides are single-stranded. The entire CpG oligodeoxynucleotide can be unmethylated or portions may be

25   unmethylated. In one embodiment, at least the C of the 5' CG 3' is unmethylated.

       Cytokine: Proteins made by cells that affect the behavior of other cells, such as lymphocytes. In one embodiment, a cytokine is a chemokine, a molecule that affects cellular trafficking.

       Epitope: An antigenic determinant. These are particular chemical groups or

30   peptide sequences on a molecule that are antigenic, *i.e.,* that elicit a specific immune response. An antibody binds a particular antigenic epitope.

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Functionally Equivalent: Sequence alterations, for example in a suppressive ODN, that yield the same results as described herein. Such sequence alterations can include, but are not limited to, deletions, base modifications, mutations, labeling, and insertions.

5   **Graft-versus-host disease:** Tissue rejection, also called graft-versus-host disease, is a consequence of organ or tissue transplantation caused by the transplant recipient's (host's) immune response to the transplanted organ/tissue which can damage or destroy it. Ordinarily, the immune response protects the body from potentially harmful substances (antigens) such as microorganisms, toxins, and

10   cancer cells. The immune system distinguishes "self" from "foreign" by reacting to proteins on the surfaces of cells. It reacts against substances it recognizes as foreign (antigens). The presence of foreign blood or tissue in the body triggers an immune response that can result in blood transfusion reactions and transplant rejection when antibodies are formed against foreign antigens on the transplanted or transfused

15   material.

       G-tetrad: G-tetrads are G-rich DNA segments that can accommodate complex secondary and/or tertiary structures (see Fig. 1). A G-tetrad involves the planar association of four Gs in a cyclic Hoogsteen hydrogen bonding arrangement (this involves non-Watson Crick base-pairing). In general, either a run of two or

20   more contiguous Gs or a hexameric region in which >50% of the bases are Gs, is needed for an ODN to form a G-tetrad. The longer the run of contiguous Gs, and the higher the G content of the ODN, the higher the likelihood of G-tetrad formation, as reflected by higher CD or ellipticity values.

       Oligonucleotides that form G-tetrads can also form higher-level aggregates

25   that are more easily recognized and taken up by immune cells, for example, through scavenger receptors or by nucleolin.

       Guanosine-rich sequence: A hexameric region of a nucleotide sequence in which >50% of the bases are Gs.

       Immunostimulatory CpG ODN: An oligodeoxynucleotide, which contains

30   a cytosine, guanine dinucleotide sequence and stimulates (*e.g.*, has a mitogenic effect) vertebrate immune cells. The cytosine, guanine is unmethylated. Both D and

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K type CpG ODNs are immunostimulatory (see in Verthelyi et al., *J. Immunol.*
166:2372-2377, 2001, which is herein incorporated by reference).

Immunosuppressive agent: A molecule, such as a chemical compound,
small molecule, steroid, nucleic acid molecule, or other biological agent, that can
5   decrease an immune response such as an inflammatory reaction.
Immunosuppressive agents include, but are not limited to an agent of use in treating
arthritis (anti-arthritis agent). Specific, non-limiting examples of
immunosuppressive agents are non-steroidal anti-inflammatory agents, cyclosporine
A, FK506, and anti-CD4. In additional examples, the agent is a biological response
10   modifier, such as Kineret® (anakinra), Enbrel® (etanercept), or Remicade®
(infliximab), a disease-modifying antirheumatic drug (DMARD), such as Arava®
(leflunomide), a nonsteroidal anti-inflammatory drug (NSAIDs), specifically a
Cyclo-Oxygenase-2 (COX-2) inhibitor, such as Celebrex® (celecoxib) and Vioxx®
(rofecoxib), or another product, such as Hyalgan® (hyaluronan) and Synvisc®
15   (hylan G-F20).

Immune-mediated disorder: A disorder that involves an unwanted
immune response. Although immune recognition of "non-self" proteins is essential
to avoid and eliminate infection, the immune response can sometimes be unwanted.
Autoimmune diseases, such as rheumatoid arthritis, multiple sclerosis or insulin
20   dependent diabetes mellitus, are the result of a pathological immune response
against self antigens, and T cells are the primary mediators of autoimmunity. For
example, rheumatoid arthritis is an autoimmune disorder, as are Hashimoto's
thyroiditis, pernicious anemia, Addison's disease, type I diabetes, systemic lupus
erythematosus, dermatomyositis, Sjogren's syndrome, dermatomyositis, lupus
25   erythematosus, multiple sclerosis, myasthenia gravis, Reiter's syndrome, and
Grave.s disease, among others.

Rejection of transplanted organs and tissues are a further example of an
immune-mediated disorder, and can often result in damage to and/or rejection of the
transplant. Tissue rejection, also called graft-versus-host disease, is a consequence
30   of organ or tissue transplantation caused by the transplant recipient's (host's)
immune response to the transplanted organ/tissue which can damage or destroy it.
Ordinarily, the immune response protects the body from potentially harmful

substances (antigens) such as microorganisms, toxins, and cancer cells. The immune system distinguishes "self" from "foreign" by reacting to proteins on the surfaces of cells. It reacts against substances it recognizes as foreign (antigens). The presence of foreign blood or tissue in the body triggers an immune response that

5 can result in blood transfusion reactions and transplant rejection when antibodies are formed against foreign antigens on the transplanted or transfused material. Before transplant, tissue is "typed" according to the antigens it contains (Histocompatibility antigens).

No two people (except identical twins) have identical tissue antigens.

10 Therefore, in the absence of immunosuppressive drugs, organ and tissue transplantation would almost always causes an immune response against the foreign tissue (rejection), which would result in destruction of the transplant. Though tissue typing ensures that the organ or tissue is as similar as possible to the tissues of the recipient, unless the donor is an identical twin, no match is perfect and the

15 possibility of organ/tissue rejection remains. Immunosuppressive therapy is used to prevent organ rejection.

Allergy is another example of an immune-mediated disorder. An allergy is a collection of symptoms caused by an exaggerated immune response or reaction to substances that do not trigger an immune response in most people. The term

20 "allergy" has become synonymous with Type I hypersensitivity (IgE-mediated allergy). Four different types of hypersensitivity were described by Coomb and Gell (Types I, II, III and IV), as a pedagogical way to increase the understanding of different immune reactions which could be provoked by many antigens. In practice these types do not necessarily occur in isolation from each other.

25 Allergic diseases generally begin in childhood, although they can arise at any age. Development of allergic disease is associated with an allergic constitution due to heredity and to environmental and health factors. An allergic response involves an increased production of allergen-specific IgE antibodies, which may lead to clinical symptoms such as rhinitis, asthma, eczema, colic pains or diarrhea. A state

30 of hyperreactivity often accompanies an allergic reaction. If this hyperreactivity occurs in the respiratory tract, everyday stimuli like dust, tobacco smoke, cold air and perfumes may lead to allergy-like symptoms.

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Immune response: A response of a cell of the immune system, such as a B cell, T cell to a stimulus. In one embodiment, the response is specific for a particular antigen (an "antigen-specific response").

Inflammation: When damage to tissue occurs, the body's response to the damage is usually inflammation. The damage may be due to trauma, lack of blood supply, hemorrhage, autoimmune attack, transplanted exogenous tissue or infection. This generalized response by the body includes the release of many components of the immune system (e.g., IL-1 and TNF), attraction of cells to the site of the damage, swelling of tissue due to the release of fluid and other processes.

Inflammatory arthropathy: An inflammatory arthropathy is an inflammatory disease affecting one or more joints, for example an inflammatory disease that affects the synovial membranes of one or more joints. Inflammatory arthropathies include, for example, arthritis, ankylosing spondylitis, Reiter's syndrome, psoriatic arthropathy, enteropathis spondylitis, juvenile arthropathy, and reactive arthropathy.

Infectious agent: An agent that can infect a subject, including, but not limited to, viruses, bacteria, and fungi.

Examples of infectious virus include: *Retroviridae* (for example, human immunodeficiency viruses, such as HIV-1 (also referred to as HTLV-III, LAV or HTLV-III/LAV, or HIV-III; and other isolates, such as HIV-LP; *Picornaviridae* (for example, polio viruses, hepatitis A virus; enteroviruses, human coxsackie viruses, rhinoviruses, echoviruses); *Calciviridae* (such as strains that cause gastroenteritis); *Togaviridae* (for example, equine encephalitis viruses, rubella viruses); *Flaviridae* (for example, dengue viruses, encephalitis viruses, yellow fever viruses); *Coronaviridae* (for example, coronaviruses); *Rhabdoviridae* (for example, vesicular stomatitis viruses, rabies viruses); *Filoviridae* (for example, ebola viruses); *Paramyxoviridae* (for example, parainfluenza viruses, mumps virus, measles virus, respiratory syncytial virus); *Orthomyxoviridae* (for example, influenza viruses); *Bungaviridae* (for example, Hantaan viruses, bunga viruses, phleboviruses and Nairo viruses); *Arena viridae* (hemorrhagic fever viruses); *Reoviridae* (e.g., reoviruses, orbiviurses and rotaviruses); *Birnaviridae*; *Hepadnaviridae* (Hepatitis B virus); *Parvoviridae* (parvoviruses); *Papovaviridae* (papilloma viruses, polyoma

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viruses); *Adenoviridae* (most adenoviruses); *Herpesviridae* (herpes simplex virus (HSV) 1 and HSV-2, varicella zoster virus, cytomegalovirus (CMV), herpes viruses); *Poxviridae* (variola viruses, vaccinia viruses, pox viruses); and *Iridoviridae* (such as African swine fever virus); and unclassified viruses (for example, the

5   etiological agents of Spongiform encephalopathies, the agent of delta hepatitis (thought to be a defective satellite of hepatitis B virus), the agents of non-A, non-B hepatitis (class 1=internally transmitted; class 2=parenterally transmitted (*i.e.,* Hepatitis C); Norwalk and related viruses, and astroviruses).

Examples of infectious bacteria include: *Helicobacter pyloris, Borelia*

10   *burgdorferi, Legionella pneumophilia, Mycobacteria sps* (such as. *M. tuberculosis, M. avium, M. intracellulare, M. kansaii, M. gordonae), Staphylococcus aureus, Neisseria gonorrhoeae, Neisseria meningitidis, Listeria monocytogenes, Streptococcus pyogenes* (Group A *Streptococcus), Streptococcus agalactiae* (Group B *Streptococcus), Streptococcus* (*viridans* group), *Streptococcus faecalis,*

15   *Streptococcus bovis, Streptococcus (anaerobic sps.), Streptococcus pneumoniae, pathogenic Campylobacter sp., Enterococcus sp., Haemophilus influenzae, Bacillus antracis, corynebacterium diphtheriae, corynebacterium sp., Erysipelothrix rhusiopathiae, Clostridium perfringers, Clostridium tetani, Enterobacter aerogenes, Klebsiella pneumoniae, Pasturella multocida, Bacteroides sp., Fusobacterium*

20   *nucleatum, Streptobacillus moniliformis, Treponema pallidium, Treponema pertenue, Leptospira,* and *Actinomyces israelli.*

Examples of infectious fungi include, but are not limited to, *Cryptococcus neoformans, Histoplasma capsulatum, Coccidioides immitis, Blastomyces dermatitidis, Chlamydia trachomatis, Candida albicans.*

25   Other infectious organisms (such as protists) include: *Plasmodium falciparum* and *Toxoplasma gondii.*

Isolated: An "isolated" biological component (such as a nucleic acid, peptide or protein) has been substantially separated, produced apart from, or purified away from other biological components in the cell of the organism in which the

30   component naturally occurs, *i.e.,* other chromosomal and extrachromosomal DNA and RNA, and proteins. Nucleic acids, peptides and proteins which have been "isolated" thus include nucleic acids and proteins purified by standard purification

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methods. The term also embraces nucleic acids, peptides and proteins prepared by recombinant expression in a host cell as well as chemically synthesized nucleic acids.

5   **Leukocyte:** Cells in the blood, also termed "white cells," that are involved in defending the body against infective organisms and foreign substances. Leukocytes are produced in the bone marrow. There are 5 main types of white blood cell, subdivided between 2 main groups: polymorphonuclear leukocytes (neutrophils, eosinophils, basophils) and mononuclear leukocytes (monocytes and lymphocytes). When an infection is present, the production of leukocytes increases.

10   **Mammal:** This term includes both human and non-human mammals. Similarly, the term "subject" includes both human and veterinary subjects.

Maturation: The process in which an immature cell, such as dendritic cell, changes in form or function to become a functional mature cell, such as an APC.

Nucleic acid: A deoxyribonucleotide or ribonucleotide polymer in either

15   single or double stranded form, and unless otherwise limited, encompasses known analogues of natural nucleotides that hybridize to nucleic acids in a manner similar to naturally occurring nucleotides.

Oligonucleotide or "oligo": Multiple nucleotides (*i.e.*, molecules comprising a sugar (*e.g.*, ribose or deoxyribose) linked to a phosphate group and to

20   an exchangeable organic base, which is either a substituted pyrimidine (Py) (*e.g.*, cytosine (C), thymine (T) or uracil (U)) or a substituted purine (Pu) (*e.g.*, adenine (A) or guanine (G)). The term "oligonucleotide" as used herein refers to both oligoribonucleotides (ORNs) and oligodeoxyribonucleotides (ODNs). The term "oligonucleotide" also includes oligonucleosides (*i.e.*, an oligonucleotide minus the

25   phosphate) and any other organic base polymer. Oligonucleotides can be obtained from existing nucleic acid sources (*e.g.*, genomic or cDNA), but are preferably synthetic (*e.g.*, produced by oligonucleotide synthesis).

A "stabilized oligonucleotide" is an oligonucleotide that is relatively resistant to *in vivo* degradation (for example via an exo- or endo-nuclease). In one

30   embodiment, a stabilized oligonucleotide has a modified phosphate backbone. One specific, non-limiting example of a stabilized oligonucleotide has a phophorothioate

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modified phosphate backbone (wherein at least one of the phosphate oxygens is replaced by sulfur). Other stabilized oligonucleotides include: nonionic DNA analogs, such as alkyl- and aryl- phophonates (in which the charged phosphonate oxygen is replaced by an alkyl or aryl group), phophodiester and

5   alkylphosphotriesters, in which the charged oxygen moiety is alkylated. Oligonucleotides which contain a diol, such as tetraethyleneglycol or hexaethyleneglycol, at either or both termini have also been shown to be substantially resistant to nuclease degradation.

An "immunostimulatory oligonucleotide," "immunostimulatory CpG

10   containing oligodeoxynucleotide," "CpG ODN," refers to an oligodeoxynucleotide, which contains a cytosine, guanine dinucleotide sequence and stimulates (*e.g.,* has a mitogenic effect) vertebrate immune cells. The cytosine, guanine is unmethylated.

An "oligonucleotide delivery complex" is an oligonucleotide associated with (*e.g.,* ionically or covalently bound to; or encapsulated within) a targeting means

15   (*e.g.,* a molecule that results in a higher affinity binding to a target cell (*e.g.,.* B-cell or natural killer (NK) cell) surface and/or increased cellular uptake by target cells). Examples of oligonucleotide delivery complexes include oligonucleotides associated with: a sterol (*e.g.,* cholesterol), a lipid (*e.g.,* cationic lipid, virosome or liposome), or a target cell specific binding agent (*e.g.,* a ligand recognized by a target cell

20   specific receptor). Preferred complexes must be sufficiently stable in vivo to prevent significant uncoupling prior to internalization by the target cell. However, the complex should be cleavable or otherwise accessible under appropriate conditions within the cell so that the oligonucleotide is functional. (Gursel, *J. Immunol.* 167: 3324, 2001)

25   **Parenteral:** Administered outside of the intestine, *e.g.,* not via the alimentary tract. Generally, parenteral formulations are those that will be administered through any possible mode except ingestion. This term especially refers to injections, whether administered intravenously, intrathecally, intramuscularly, intraperitoneally, intra-articularly, or subcutaneously, and various

30   surface applications including intranasal, intradermal, and topical application, for instance.

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Pharmaceutical agent or drug: A chemical compound or composition capable of inducing a desired therapeutic or prophylactic effect when properly administered to a subject. Pharmaceutical agents include, but are not limited to, chemotherapeutic agents and anti-infective agents.

5   **Pharmaceutically acceptable carriers:** The pharmaceutically acceptable carriers useful in this disclosure are conventional. *Remington's Pharmaceutical Sciences*, by E. W. Martin, Mack Publishing Co., Easton, PA, 15th Edition (1975), describes compositions and formulations suitable for pharmaceutical delivery of the fusion proteins herein disclosed.

10   In general, the nature of the carrier will depend on the particular mode of administration being employed. For instance, parenteral formulations usually comprise injectable fluids that include pharmaceutically and physiologically acceptable fluids such as water, physiological saline, balanced salt solutions, aqueous dextrose, glycerol or the like as a vehicle. For solid compositions (*e.g.*,

15   powder, pill, tablet, or capsule forms), conventional non-toxic solid carriers can include, for example, pharmaceutical grades of mannitol, lactose, starch, or magnesium stearate. In addition to biologically-neutral carriers, pharmaceutical compositions to be administered can contain minor amounts of non-toxic auxiliary substances, such as wetting or emulsifying agents, preservatives, and pH buffering

20   agents and the like, for example sodium acetate or sorbitan monolaurate.

Preventing or treating a disease: "Preventing" a disease refers to inhibiting the full development of a disease, for example in a person who is known to have a predisposition to a disease such as an autoimmune disorder. An example of a person with a known predisposition is someone with a history of diabetes in the

25   family, or who has been exposed to factors that predispose the subject to a condition, such as lupus or arthritis. "Treatment" refers to a therapeutic intervention that ameliorates a sign or symptom of a disease or pathological condition after it has begun to develop.

Purified: The term purified does not require absolute purity; rather, it is

30   intended as a relative term. Thus, for example, a purified peptide preparation is one in which the peptide or protein is more enriched than the peptide or protein is in its natural environment within a cell. Preferably, a preparation is purified such that the

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protein or peptide represents at least 50% of the total peptide or protein content of the preparation.

Suppressive ODN: DNA molecules of at least eight nucleotides in length, wherein the oligodeoxynucleotide forms a G-tetrad, and has a CD value of greater

5   than about 2.9. In a suppressive ODN the number of guanosines is at least two. In one embodiment, a suppressive ODN inhibits immune activation caused by CpG DNA when administered prior to, concurrently with, or after the administration of an CpG ODN. at least about 8 nucleotides in length.

Therapeutic agent: Used in a generic sense, it includes treating agents,

10   prophylactic agents, and replacement agents.

Therapeutically effective amount of [a compound]: A quantity of a specified compound sufficient to achieve a desired effect in a subject being treated. For instance, this can be the amount of a suppressive ODN necessary to suppress CpG-induced immune cell activation in a subject, or a dose sufficient to prevent

15   advancement, or to cause regression of a disease, or which is capable of relieving symptoms caused by a disease, such as pain or swelling..

An effective amount of a suppressive ODN can be administered systemically or locally. In addition, an effective amount of a suppressive ODN can be administered in a single dose, or in several doses, for example daily, during a course

20   of treatment. However, the effective amount of the ODN will be dependent on the preparation applied, the subject being treated, the severity and type of the affliction, and the manner of administration of the compound. For example, a therapeutically effective amount of a suppressive ODN can vary from about 0.01 mg/kg body weight to about 1 g/kg body weight in some embodiments, or from about 0.01

25   mg/kg to about 60 mg/kg of body weight, based on efficacy.

The suppressive ODNs disclosed herein have equal applications in medical and veterinary settings. Therefore, the general terms "subject" and "subject being treated" are understood to include all animals, including humans or other simians, dogs, cats, horses, and cows.

30   **Therapeutically effective dose:** A dose sufficient to prevent advancement, or to cause regression of the disease, or which is capable of relieving symptoms caused by the disease, such as pain or swelling.

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III.   *Description of Several Embodiments*

A.   *Suppressive Oligodeoxynucleotides and Guanosine-quadruplexes (G-tetrads)*

The present disclosure relates to a class of DNA motifs that selectively

5   inhibits or suppresses immune activation. Optimal activity is observed using

multimers of these motifs, which are rich in G bases and capable of forming G-

quadruplexes (G-tetrads). G-tetrads are G-rich DNA segments that can

accommodate complex secondary and/or tertiary structures (see Fig. 1). The

suppressive ODNs of the disclosure are highly specific (*i.e.,* are neither toxic nor

10   non-specifically immunosuppressive), and are useful for inhibiting an immune

response. In one embodiment, a suppressive ODN is of use for blocking

immunostimulation caused by CpG motifs *in vivo* and *in vitro*.

A G-tetrad involves the planar association of four Gs in a cyclic Hoogsteen

hydrogen bonding arrangement (this involves non-Watson Crick base-pairing). In

15   general, either a run of two or more contiguous Gs or a hexameric region in which

>50% of the bases are Gs, is needed for an ODN to form a G-tetrad. The longer the

run of continuous Gs, and the higher the G content of the ODN, the higher the

likelihood of G-tetrad formation, as reflected by higher ellipticity values.

Oligonucleotides that form G-tetrads can also form higher-level aggregates that are

20   more easily recognized and taken up by immune cells, for example, through

scavenger receptors or by nucleolin.

The formation of G-tetrads yields a complex with different physical

properties than the individual oligonucleotides. Spectroscopically, this is manifested

by an increase in circular dicroism (CD), and an increase in peak absorbance to the

25   260-280 nm wavelength owing to the formation of secondary structures. Thus, a

convenient method for identifying oligonucleotides that form G-tetrads is to study

their CD values. An increase in peak ellipticity values to greater than 2.0 is typical

of a G-tetrad forming oligonucleotide. For instance, G-tetrad-forming ODNs can

have CD values of 2.2, 2.4, 2.6, 2.8, 3.0, 3.2, 3.5, 4.0, 4.5, 5.0, 5.5, 6.0, 6.5, 7.0, 7.5,

30   8.0, 8.5, 9.0, or higher. The higher the ellipticity value, the greater the tetrad-

forming capacity of the oligonucleotide, so an ODN with a CD value of 8.5 is

typically more suppressive than an ODN with a CD value of 2.9.

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In some embodiments, the ODN is from about 8 to about 120 nucleotides in length. In particular examples, the ODN is from about 10 to about 30 nucleotides in length. Optionally, the suppressive ODN has multiple guanosine-rich sequences, for example, in certain embodiments the ODN has from about two to about 20

5   guanosine-rich sequences, or, more particularly, from about two to about four guanosine-rich sequences.

In one embodiment, the suppressive ODNs have a sequence comprising at least one of the human telomere-derived TTAGGG suppressive motifs (see Example 1). In some examples, the ODN has at least one TTAGGG motif, and in certain

10   examples, the ODN has multiple TTAGGG motifs. For example, in particular examples, the ODN has from about two to about 20 TTAGGG motifs, or from about two to about four TTAGGG motifs. In this embodiment, suppressive ODNs containing multiple TTAGGG repeats are the most suppressive. Single TTAGGG motifs are suppressive only when incorporated into larger ODNs with greater than

15   10 bases. The TTAGGG motifs may be in either the *cis* or *trans* position, *i.e.*, they may be present on the same or on a different strand of DNA than that expressing the stimulatory CpG sequence.

Suppression of CpG-induced immune activation requires a G-tetrad-forming sequence that imposes the two-dimensional structure necessary for G-tetrad

20   formation. Examples of suppressive ODN include, but are not limited to, those shown in Fig. 9. However, any oligonucleotide capable of forming G-tetrads may be used to suppress CpG DNA-induced immune activation. In particular examples, the ODN has a sequence selected from the group consisting of SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID

25   NO: 7, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO: 10, SEQ ID NO: 11, SEQ ID NO: 12, SEQ ID NO: 13, SEQ ID NO: 14, SEQ ID NO: 15, SEQ ID NO: 16, SEQ ID NO: 17, SEQ ID NO: 18, SEQ ID NO: 19, SEQ ID NO: 20, SEQ ID NO: 21, SEQ ID NO: 22, SEQ ID NO: 23, SEQ ID NO: 24, and SEQ ID NO: 25.

Furthermore, in particular embodiments the ODN is modified to prevent

30   degradation. In one embodiment, suppressive ODNs can include modified nucleotides to confer resistance to degradation. Without being bound by theory, modified nucleotides can be included to increase the stability of a suppressive ODN.

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Thus, because phosphorothioate-modified nucleotides confer resistance to exonuclease digestion, the suppressive ODNs are "stabilized" by incorporating phosphorothioate-modified nucleotides.

In some embodiments, the ODN has a phosphate backbone modification, and in particular examples, the phosphate backbone modification is a phosphorothioate backbone modification. In one embodiment, the guanosine-rich sequence and its immediate flanking regions include phosphodiester rather than phosphorothioate nucleotides. In one specific non-limiting example, the sequence TTAGGG includes phosphodiester bases. In some examples, all of the bases in an ODN are phosphodiester bases. In other examples, the ODN is a phosphorothioate/phosphodiester chimera.

As disclosed herein, any suitable modification can be used to render the ODN resistant to degradation *in vivo* (*e.g.*, via an exo- or endo-nuclease). In one specific, non-limiting example, a modification that renders the ODN less susceptible to degradation is the inclusion of nontraditional bases such as inosine and quesine, as well as acetyl-, thio- and similarly modified forms of adenine, cytidine, guanine, thymine, and uridine. Other modified nucleotides include nonionic DNA analogs, such as alkyl or aryl phosphonates (*i.e.*, the charged phosphonate oxygen is replaced with an alkyl or aryl group, as set forth in U.S. Patent No. 4,469,863), phosphodiesters and alkylphosphotriesters (*i.e.*, the charged oxygen moiety is alkylated, as set forth in U.S. Patent No. 5,023,243 and European Patent No. 0 092 574). ODNs containing a diol, such as tetraethyleneglycol or hexaethyleneglycol, at either or both termini, have also been shown to be more resistant to degradation.

The suppressive ODN of the disclosure can be synthesized by standard methods well known in the art. Most commonly, synthesis is performed on an oligonucleotide synthesizer using the standard cyanoethyl phosphoramidite chemistry. These include, but are not limited to, phosphodiester, phosphorothioate, peptide nucleic acids, synthetic peptide analogues, and any combination thereof. Those skilled in the art will recognize that any other standard technique may be used to synthesize the suppressive ODN described herein.

In one embodiment, a suppressive ODN is included in a delivery complex. The delivery complex can include the suppressive ODN and a targeting means. Any

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suitable targeting means can be used. For example, in some embodiments, a suppressive ODN is associated with (*e.g.*, ionically or covalently bound to, or encapsulated within) a targeting means (*e.g.*, a molecule that results in higher affinity binding to a target cell, such as a B cell). A variety of coupling or cross-

5   linking agents can be used to form the delivery complex, such as protein A, carbodiamide, and N-succinimidyl-3-(2-pyridyldithio) propionate (SPDP). Examples of oligodeoxynucleotide delivery complexes include a suppressive ODN associated with a sterol (*e.g.*, cholesterol), a lipid (*e.g.*, a cationic lipid, anionic lipid, virosome or liposome), and a target cell specific binding agent (*e.g.*, a ligand

10   recognized by target cell specific receptor). Without being bound by theory, the complex is sufficiently stable *in vivo* to prevent significant uncoupling prior to delivery to the target cell. In one embodiment, the delivery complex is cleavable such that the ODN is released in a functional form at the target cells.

15   *B.   Pharmaceutical Compositions*

The suppressive ODNs described herein may be formulated in a variety of ways depending on the location and type of disease to be treated. Pharmaceutical compositions are thus provided for both local (*e.g.* topical or intra-articular) use as well as for systemic use. Therefore, the disclosure includes within its scope

20   pharmaceutical compositions comprising at least one suppressive ODN formulated for use in human or veterinary medicine.

Pharmaceutical compositions that include at least one suppressive ODN as described herein as an active ingredient, or that include both a suppressive ODN and an additional anti-inflammatory or anti-arthritis factor as active ingredients, may be

25   formulated with an appropriate solid or liquid carrier, depending upon the particular mode of administration chosen. Additional active ingredients include, for example, non-steroidal anti-inflammatory agents, such as diclofenac, diflunisal, etodolac, flurbiprofen, ibuprofen, indomethacin, ketoprofen, ketorolac, nabumetone, naproxen, oxaprozin, piroxicam, sulindac, tolmetin, celecoxib, and rofecoxib,

30   steroids, such as cortisone, dexamethasone, hydrocortisone, methylprednisolone,

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prednisolone, prednisone, and triamcinolone, and immunosuppressives, for example cyclosporin, tacrolimus, mycophenolic acid, and sirolimus.

The pharmaceutically acceptable carriers and excipients useful in this disclosure are conventional. For instance, parenteral formulations usually comprise

5   injectable fluids that are pharmaceutically and physiologically acceptable fluid vehicles such as water, physiological saline, other balanced salt solutions, aqueous dextrose, glycerol or the like. Excipients that can be included are, for instance, proteins, such as human serum albumin or plasma preparations. If desired, the pharmaceutical composition to be administered may also contain minor amounts of

10   non-toxic auxiliary substances, such as wetting or emulsifying agents, preservatives, and pH buffering agents and the like, for example sodium acetate or sorbitan monolaurate.

The dosage form of the pharmaceutical composition will be determined by the mode of administration chosen. For instance, in addition to injectable fluids,

15   topical and oral formulations can be employed. Topical preparations can include eye drops, ointments, sprays and the like. Oral formulations may be liquid (*e.g.*, syrups, solutions, or suspensions), or solid (*e.g.*, powders, pills, tablets, or capsules). For solid compositions, conventional non-toxic solid carriers can include pharmaceutical grades of mannitol, lactose, starch, or magnesium stearate. Actual

20   methods of preparing such dosage forms are known, or will be apparent, to those of ordinary skill in the art.

The pharmaceutical compositions that comprise a suppressive ODN, in some embodiments, will be formulated in unit dosage form, suitable for individual administration of precise dosages. The amount of active compound(s) administered

25   will be dependent on the subject being treated, the severity of the affliction, and the manner of administration, and is best left to the judgment of the prescribing clinician. Within these bounds, the formulation to be administered will contain a quantity of the active component(s) in amounts effective to achieve the desired effect in the subject being treated.

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C.   Therapeutic Uses

A method is disclosed herein for treating or preventing an immune mediated disorder in a subject. In one specific, non-limiting example, the immune mediated disorder is an autoimmune disease. In another specific, non-limiting example, the immune mediated disorder is an allergic reaction. In a further specific, non-limiting example, the immune mediated disorder is transplant rejection.

Autoimmune diseases include, but are not limited to inflammatory arthritis, Hashimoto's thyroiditis, pernicious anemia, Addison's disease, type I diabetes, systemic lupus erythematosus, dermatomyositis, Sjogren's syndrome, dermatomyositis, lupus erythematosus, multiple sclerosis, myasthenia gravis, Reiter's syndrome, and Grave.s disease, among others.

Rejection of transplanted organs and tissues are a further example of an undesired consequence of normal immunity, which can often result in damage to and/or rejection of the transplant. Tissue rejection, also called graft-versus-host disease, is a consequence of organ or tissue transplantation caused by the transplant recipient's (host's) immune response to the transplanted organ/tissue which can damage or destroy it. Ordinarily, the immune response protects the body from potentially harmful substances (antigens) such as microorganisms, toxins, and cancer cells. The immune system distinguishes "self" from "foreign" by reacting to proteins on the surfaces of cells. It reacts against substances it recognizes as foreign (antigens). The presence of foreign blood or tissue in the body triggers an immune response that can result in blood transfusion reactions and transplant rejection when antibodies are formed against foreign antigens on the transplanted or transfused material. Before transplant, tissue is "typed" according to the antigens it contains (Histocompatibility antigens).

No two people (except identical twins) have identical tissue antigens. Therefore, in the absence of immunosuppressive drugs, organ and tissue transplantation would almost always causes an immune response against the foreign tissue (rejection), which would result in destruction of the transplant. Though tissue typing ensures that the organ or tissue is as similar as possible to the tissues of the recipient, unless the donor is an identical twin, no match is perfect and the

possibility of organ/tissue rejection remains. Immunosuppressive therapy is used to prevent organ rejection.

Allergy is another example of an immune-mediated disorder. An allergy is a collection of symptoms caused by an exaggerated immune response or reaction to

5   substances that do not trigger an immune response in most people. The term "allergy" has become synonymous with Type I hypersensitivity (IgE-mediated allergy). Four different types of hypersensitivity were described by Coomb and Gell (Types I, II, III and IV), as a pedagogical way to increase the understanding of different immune reactions which could be provoked by many antigens. In practice

10   these types do not necessarily occur in isolation from each other.

Allergic diseases generally begin in childhood, although they can arise at any age. Development of allergic disease is associated with an allergic constitution due to heredity and to environmental and health factors. An allergic response involves an increased production of allergen-specific IgE antibodies, which may lead to

15   clinical symptoms such as rhinitis, asthma, eczema, colic pains or diarrhea. A state of hyperreactivity often accompanies an allergic reaction. If this hyperreactivity occurs in the respiratory tract, everyday stimuli like dust, tobacco smoke, cold air and perfumes may lead to allergy-like symptoms.

The method includes administering a therapeutically effective amount of the

20   suppressive ODN to a subject having or at risk of developing an immune-mediated disorder, thereby treating or preventing the immune-mediated disorder. In one embodiment, the suppressive ODN can be administered locally, such as by intra-articular injection. In another embodiment, the suppressive ODN is administered systemically. In one embodiment, the immune-mediated disorder is an autoimmune

25   disease, an allergy, or graft-versus-host disease. In particular embodiments, the immune-mediated disorder is an inflammatory arthropathy.

In order to treat or prevent an immune-mediated disorder, a therapeutically effective amount of a suppressive ODN (see above) is administered to the subject. In one embodiment, the ODN has a CD value of greater than about 2.9, 3.0, 3.2, 3.4,

30   3.6, 3.8, 4.0, or 5.0. In some embodiments of the method, the ODN is from about 8 to about 120 nucleotides in length. In particular examples of the method, the ODN is from about 10 to about 30 nucleotides in length. Optionally, the suppressive ODN

has multiple guanosine-rich sequences, for example, in certain embodiments of the method, the ODN has from about two to about 20 guanosine-rich sequences, or, more particularly, from about two to about four guanosine-rich sequences.

In some examples of the method, the ODN has at least one TTAGGG motif, and in certain examples, the ODN has multiple TTAGGG motifs. For example, in particular examples of the method, the ODN has from about two to about 20 TTAGGG motifs, or from about two to about four TTAGGG motifs. In particular examples of the method, the ODN has a sequence selected from the group consisting of SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 7, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO: 10, SEQ ID NO: 11, SEQ ID NO: 12, SEQ ID NO: 13, SEQ ID NO: 14, SEQ ID NO: 15, SEQ ID NO: 16, SEQ ID NO: 17, SEQ ID NO: 18, SEQ ID NO: 19, SEQ ID NO: 20, SEQ ID NO: 21, SEQ ID NO: 22, SEQ ID NO: 23, SEQ ID NO: 24, and SEQ ID NO: 25.

Combinations of these suppressive ODN are also of use. Thus, in one embodiment, more than one suppressive ODN, each with a different nucleic acids sequence, are administered to the subject. In several specific, non-limiting examples, at least two, at least three, or at least four suppressive ODN are administered to the subject.

In another embodiment, an additional anti-inflammatory agent or immunosuppressive agent is administered in conjunction with a suppressive ODN. The administration of the anti-inflammatory agent or immunosuppressive agent and the suppressive ODN can be sequential or simultaneous.

In particular examples, the immunosuppressive agent is a non-steroidal anti-inflammatory agent, such as diclofenac, diflunisal, etodolac, flurbiprofen, ibuprofen, indomethacin, ketoprofen, ketorolac, nabumetone, naproxen, oxaprozin, piroxicam, sulindac, tolmetin, celecoxib, or rofecoxib, a steroid, such as cortisone, dexamethasone, hydrocortisone, methylprednisolone, prednisolone, prednisone, or triamcinolone, or an immunosuppressive agent, for example cyclosporin, tacrolimus, mycophenolic acid, or sirolimus.

In another embodiment an additional anti-inflammatory agent is administered in conjunction with a suppressive ODN. The agent can be an

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immunosuppressive, or an anti-arthritis agent. The administration of the anti-inflammatory agent and the suppressive ODN can be sequential or simultaneous.

In particular examples, the immunosuppressive agent is a biological response modifier, such as Kineret® (anakinra), Enbrel® (etanercept), or Remicade®
.5   (infliximab), a disease-modifying antirheumatic drug (DMARD), such as Arava® (leflunomide), a nonsteroidal anti-inflammatory drug (NSAIDs), specifically a Cyclo-Oxygenase-2 (COX-2) inhibitor, such as Celebrex® (celecoxib) and Vioxx® (rofecoxib), or another product, such as Hyalgan® (hyaluronan) and Synvisc® (hylan G-F20).

10   Thus, the suppressive ODNs disclosed herein may be administered to a subject for the treatment of immune-mediated disorders in that individual. ODN administration can be systemic or local. Local administration of the ODN is performed by methods well known to those skilled in the art. By way of example, one method of administration to the knee, hip and/or shoulder of an individual is by
15   intra-articular injection. For administration to the knee, for example, the joint to be injected is washed with a betadine solution or other antiseptic. A solution of about one percent lidocaine hydrochloride is injected into the skin and subcutaneous tissue. A 3-way stopcock/needle assembly is utilized to administer the compound via an 18-30 gauge needle. The ODN is injected into the joint space using a
20   standard lateral approach well known to those skilled in the art. The needle and needle tract are cleansed by flushing with 1% lidocaine hydrochloride through the 3-way stopcock assembly as the needle is withdrawn. The knee is then moved through a flexion-extension arc and then immobilized in full extension. The patient is then confined to bed for approximately 24 hours to minimize movement and minimize
25   leakage of ODN from the joint.

In other embodiment, the administration of the suppressive ODN is systemic. Oral, intravenous, intra-arterial, subcutaneous, intra-peritoneal, intra-muscular, and even rectal administration is contemplated.

In other embodiments, the method is a method of treating or preventing an
30   autoimmune disease in a subject that involves contacting immune cells with a suppressive ODN, and transferring the immune cells to a subject having or at risk of developing an autoimmune disease, thereby treating or preventing the autoimmune

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disease. Without being bound by theory, these immune cells act to suppress immune activation in a subject. One specific, non-limiting example is dendritic cells. Thus, in certain examples, the immune cells, such as dendritic cells, are contacted with a suppressive ODN, and subsequently administered to a subject. The

5 immune cells can be delivered alone, in conjunction with a suppressive ODN, and/or in conjunction with an additional immunosuppressive agent. The immune cells can be delivered either systemically or locally. In several specific, non-limiting examples, the cells are delivered parenterally, intravenously, intra-muscularly, sub-cutaneously, or intra-articularly.

10 Precise, effective quantities of cells can be readily determined by those who are skilled in the art and will depend, of course, upon the exact condition being treated by the particular therapy being employed. The cells can be transplanted to a desired location, or can be administered intravenously. Other agents, such as growth factors or immunosuppressive agents, can be administered in conjunction with the

15 immune cells.

In other embodiments, suppressive ODNs are used to inhibit angiogenesis. In one embodiment, suppressive ODNs are utilized to inhibit angiogenesis *in vivo*. Thus, a suppressive ODN and a pharmacologically acceptable carrier are administered to a subject, such that angiogenesis is inhibited in the subject. Suitable

20 subjects include, but are not limited to, subjects with a tumor, subjects with macular degeneration, or subjects with diabetic retinopathy. Suitable tumors include, but are not limited to bone, brain, breast, gastrointestinal, endocrine, eye, genitourinary, germ cell, gynecologic, head, neck, hematologic, leukemic, lung, lymphm, musculoskeletal, neurologic, respiratory, thoracic, and skin tumors.

25 In another embodiment, the CpG ODN is administered with a second anti-angiogenic factor. Specific, non-limiting examples of anti-angiogenic factors of use include, but are not limited to Macugen, rhuFab, alpha-interferon, thalidomide and prinomastat. The second anti-angiogenic agent can be administered simultaneously or sequentially with the suppressive ODN.

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D.   *Kits*

Further embodiments of the disclosure include kits useful for administering

the suppressive ODN described herein *in vivo* or *in vitro*. For example, a kit useful

for treating a subject with an autoimmune disease would comprise an appropriate

5   dosage of suppressive ODN, as well as, optionally, any agents useful for enhancing

the inhibitory effect of suppressive ODN. Other embodiments further include

instructions for using the kit, and/or pre-filled syringes for administering the ODN to

a subject.

Thus, in one embodiment, a kit is provided including a container of

10   suppressive ODN (a sufficient amount for either a single use or multiple uses), and

instructions the suppressive ODN. The instructions can be in written form, or can be

provided in an electronic format, such as on a diskette or a CD-ROM. Instructions

can also be provided in the form of a video cassette.

The subject matter of the present disclosure is further illustrated by the

15   following non-limiting Examples.

EXAMPLES

Example 1

Repetitive Elements Present in Mammalian Telomeres Suppress Bacterial

20   **DNA-Induced Immune Activation**

This example demonstrates the ability of TTAGGG multimers to inhibit

CpG-induced immune activation.

A.   *General Methods*

25   *Reagents*

Endotoxin-free phosphorothioate or phosphodiester ODNs were synthesized

at the CBER core facility. 7-DG modified ODNs were synthesized using the 10-

camphorsulphonyl-oxaziridine oxidization protocol recommended by the

manufacturer (Glen Research, Sterling, VA). Mammalian DNA was isolated from

30   calf thymus and murine spleen (Wizard Genomic DNA purification kit, Promega,

Madison, WI). E. coli was obtained from Sigma (St. Louis, MO). Telomerase

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knockout mice were provided by Dr. Carol Greider (Johns Hopkins Univ.,

Baltimore, MD). All DNA obtained from commercial providers was re-purified to

eliminate endotoxin (<0.1 U/mg). Double stranded DNA was converted to single

stranded DNA by heat denaturing at 95°C for 5' followed by cooling on ice. BAL-

5   31 (New England Biolabs, Beverly, MA) digestion of CT DNA was continued for

2h at 30°C according to manufacturer's recommendations. At the end of the

incubation, the enzyme was inactivated at 65°C for 10 min. Plasmid encoding for

1.6 kb long TTAGGG repeat was a generous gift from Dr. Jerry Shay (University of

Texas Southwestern Medical Center, Dallas, TX). Non-telomere coding plasmid

10   was from Vical (San Diego, CA.).

Mice

Specific pathogen-free male BALB/c mice (Jackson Laboratories, Bar

Harbor, ME) were housed in sterile micro-isolator cages in a barrier environment

15   and injected *i.p.* with 400 g of CpG ODN plus 200 g of suppressive or control ODN.

Spleen cells were harvested 6 h later and monitored for cytokine production after 36

h. In order to measure the pEPO transgene expression levels, female Balb/c mice

(4-6 weeks old) were injected with 30 µg of pVRmEPO (a kind gift of Vical, San

Diego, CA) plasmid in sterile saline into the anterior tibialis muscle alone or in

20   combination with 50 µg of control or suppressive ODN. Hematocrits were

measured as described (Tripathy *et al., Proc Natl Acad Sci U S A.* 93:10876-80,

1996) on blood collected by tail vein puncture.

Cytokine and IgM ELISA assays

25   Immulon 2 microtiter plates (Dynex Technologies Inc., Chantilly VA) were

coated with anti-cytokine or anti-IgM antibodies (Pharmingen, San Diego CA) and

then blocked with PBS - 1% BSA. Serially diluted culture supernatant or serum was

added for 2 h. Cytokine was detected using biotinylated anti-cytokine Ab followed

by phosphatase-streptavidin (Pharmingen) whereas bound IgM was detected using

30   phosphatase-conjugated anti-IgM antibodies (Southern Biotechnology Associates,

Birmingham, AL) as described.

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Detection of co-stimulatory molecule expression by FACS

2×10^6 spleen cells/ml were incubated with ODN for 24 h. Cells were washed, fixed with 5% paraformaldehyde for 15 min, and stained with PE-labeled anti CD-40, anti CD-86, and anti ICAM-1 (Pharmingen, San Diego, CA.) for 30 min

5   at RT. Cells were washed, re-suspended in PBS/BSA (supplemented with Azide), and analyzed by FACSort (Becton Dickinson, San Jose, CA).

Cytokine RT-PCR

Spleen cells were isolated 6 h after CpG ODN injection. Total RNA was

10   extracted, reverse-transcribed, and amplified in a standard PCR reaction for 24 cycles using primers specific for murine IL-6, IL-12, and IFN as previously described (Takeshita *et al., Neuroreport* (2001)). PCR amplified material was separated on 1.5 % agarose gels and visualized under UV light after ethidium bromide staining.

15

Cell transfection and luciferase assay

HEK 293 (5×10^4) cells (ATCC, Manassas, VA) were transfected with 0.8 µg of vector plasmid (pCIneo, Promega, Madison, WI), pCIneo-mTLR9, plus 0.1 µg of p5xNF- kB-luc (Stratagene, Lajolla, CA.) and 0.1 µg of pSV-beta-galactosidase

20   (Promega) and incubated overnight at 37°C. The cells were then stimulated with indicated ODN for 24 h. Cells were then harvested and luciferase assay was performed as recommended by the manufacturer (Promega). Beta-galactosidase activity was used to normalize the data.

25   *Measurement of circular dichroism*

A Jasco J-720A spectropolarimeter was used to measure the circular dichroism of ODN (50 g/ml in 0.1X PBS). Data is expressed as the mean peak ellipticity (mdeg/abs) of 5-10 readings/sample in the 260-270 nm range.

30   *Statistical analysis*

In vitro assays were performed in triplicate on at least 3 different spleen cell preparations. All in vivo experiments were performed on a minimum of 5-10

mice/group. Statistical significance was evaluated using Student's t test. Correlation analysis is computed by linear correlation analysis between CD data vs % suppression.

5   B.   *TTAGGG multimers inhibit CpG-induced immune activation*

The ability of TTAGGG multimers to inhibit CpG-induced immune activation was tested. Initial experiments showed that ODNs containing suppressive motifs inhibit CpG-induced immune activation in a dose-dependent fashion (Fig. 2A).

10   ODNs containing the largest number of TTAGGG repeats (n=4) were the most suppressive (p<0.001; Fig. 2B). Single TTAGGG hexamers were suppressive only when incorporated into larger ODNs (≥10 bases in length), and were somewhat less active than TTAGGG multimers. TTAGGG motifs were suppressive both in *cis* and in *trans, i.e.,* they inhibited immune activation when present on the same or on a

15   different strand of DNA than that expressing the stimulatory CpG sequence (Fig. 2C and D). This inhibitory activity was exquisitely specific: even high concentrations of suppressive ODNs had no effect on mitogen-induced immune responses (Fig. 2D), indicating that suppressive ODN were neither toxic nor non-specifically immunosuppressive.

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C.   *Suppression is mediated by poly-G sequences*

The bases contributing to this suppressive activity were identified by systematically modifying ODNs containing single TTAGGG motifs. Substitutions outside the telomere-derived sequence did not significantly affect suppression.

25   Replacing multiple bases in the TTA region of the TTAGGG motif also had little effect on the suppressive activity (Fig. 2E). In contrast, replacing two or more of the Gs in this motif substantially reduced the ODN's ability to block CpG-induced immune activation (Fig. 2E). These results suggest that suppression is mediated by the poly-G sequence itself, or the two-dimensional structure imposed by that

30   sequence.

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D.   *Suppression is mediated by the two-dimensional structure of the motifs*

        To differentiate between these alternatives, individual Gs were replaced by
7-deaza guanosine (7-DG) analogues. These 7-DG substitutions did not alter the
base sequence of the ODN but did prevent Hoogsteen hydrogen bonding between

5   guanosines, thereby reducing G tetrad formation (Hurley *et al., Trends Pharmacol.
Sci.* 21:136-142, 2000; Lilley *et al., EMBO J.* 13:993-1001, 1994). The resultant
loss in secondary structure is reflected by a loss in circular dichroism (Lilley *et al.,
EMBO J.* 13:993-1001, 1994). ODNs capable of forming G tetrads typically have
peak CD values >2, while those without such secondary structure have circular

10   dichroism values <1.4 (Fig. 2).

        Introducing a 7-DG substitution significantly reduced an ODN's ability to
form a G-tetrad and to mediate suppression (p <0.001, Fig. 3). To confirm that G-
tetrad formation was critical to suppression, ODNs were synthesized that lacked the
TTAGGG motif but still formed G-tetrads. These novel ODNs suppressed CpG

15   ODN and bacterial DNA induced immune stimulation by >90% (p<0.001; Fig. 2F).
Indeed, there was a consistent correlation between G-tetrad forming ability and
suppressive activity (R = 0.832, Fig. 3B).

E.   *Mechanism of suppressive ODN activity*

20   The mechanism by which suppressive ODNs inhibit immune activation was
explored. We confirmed that both bacterial DNA and CpG ODN stimulate DNA-
dependent protein kinase activity *in vitro*. Of particular interest, this activity was
specifically blocked by TTAGGG multimers and other G tetrad forming ODN, but
not their 7-DG modified analogues. These findings indicate that suppressive motifs

25   inhibit CpG-dependent DNA-PK$_{os}$ activation of immune cells. In contrast,
suppressive ODNs did not block the binding or internalization of CpG ODNs
mediated by Toll-like receptor 9.

Example 2

30   **Effect of Suppressive ODNs on CpG-induced Immune Action**

        This example demonstrates the kinetics, magnitude, and nature of the
immune inhibition elicited by suppressive motifs. Previous studies established that

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the immunostimulatory activity of CpG DNA can be reversed several hours later either by removing the stimulatory DNA or adding suppressive DNA. The same cells that interact with stimulatory motifs also recognize suppressive motifs. When both sequence types are present on the same strand of DNA, recognition proceeds in

5   a 5' —> 3' direction. Suppression is generally dominant over stimulation, although a motif in the 5' position can interfere with recognition of a motif immediately downstream. Understanding the rules governing cellular responses to stimulatory and suppressive motifs facilitates the design of ODN for therapeutic uses.

10 *A.   General Methods*
    Animals:
          Female Balb/c mice were obtained from the Jackson Laboratories (Bar Harbor, Maine). The mice were housed under specific pathogen free conditions, and used at 8-20 weeks of age. All studies involved protocols approved by the CBER

15 Animal Care and Use Committee.

    Oligodeoxynucleotides:
          Studies utilized phosphorothioate modified ODNs that were synthesized at the CBER core facility Verthelyi, D., *et al.*, *J Immunol* (2001) 166:2372. The

20 following ODNs were used: immunostimulatory ODN1466 (TCAACGTTGA; SEQ. ID NO: 26) and ODN1555 (GCTAGACGTTAGCGT; SEQ. ID NO: 27), control ODN1471 (TCAAGCTTGA; SEQ. ID NO: 28) and ODN1612 (GCTAGAGCTTAGGCT; SEQ. ID NO: 29), suppressive ODN1502 (GAGCAAGCTGGACCTTCCAT; SEQ. ID NO: 20) and ODNH154

25 (CCTCAAGCTTGAGGGG; SEQ. ID NO: 1). Underlined bases represent the 10-mer sequences that were incorporated into complex multi-determinant ODN used in some experiments. There was no detectable protein or endotoxin contamination of these ODN.
          Mammalian DNA was purified from BALB/c spleens (Wizard Genomic

30 DNA purification kit, Promega, Madison, WI). E. coli DNA was obtained from Gibco BRL (Rockville, MD, USA). Endotoxin contamination in these preparations was <0.1 U/ml after purification Klinman, D. M., *et al.*, *J.Immunol.* (1997)

158:3635. Double stranded DNA (dsDNA) was converted to single stranded DNA (ssDNA) by heat denaturing at 95°C for 5' followed by immediate cooling on ice.

Cytokine ELISA assays:

5   Single spleen cell suspensions were washed 3 times and re-suspended in RPMI- 1640 supplemented with 5% heat inactivated fetal calf serum (FCS), 1.5 mM L- glutamine and 100 U/ml of penicillin/streptomycin. 5 x 10^5 cells/well were cultured in flat-bottomed microtiter plates (Costar, Corning, New York) with 1 uM ODN for 18 - 24 h. Culture supernatants were collected, and cytokine levels

10   measured by ELISA. In brief, 96 well Immulon H2B plates were coated with cytokine-specific antibodies and blocked with PBS 1% BSA as previously described in Klinman, D.M., and Nutman, T.B., *Current Protocols in Immunology* (1994), J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M. Shevach, and W. Strober, eds., Greene Publishing Associates, Brooklyn, NY. Culture supernatants were added, and

15   bound cytokine detected by the addition of biotin labeled secondary antibodies followed by phosphatase-conjugated avidin and a phosphatase-specific colorimetric substrate (PNPP, Pierce, Rockford, Illinois). Standard curves were generated using recombinant cytokines. The detection limit for these assays was: 0.8 U/ml for IFNg, 0.1 ng/ml for IL-6 and 0.1 ng/ml for IL-12. All assays were performed in triplicate.

20

Cytokine-specific ELIspot assays:

        A single spleen cell suspension prepared in RPMI 1640 plus 5% FCS was serially diluted onto plates pre-coated with anti-cytokine Abs. Klinman, D.M., and Nutman, T.B., *Current Protocols in Immunology* (1994), J. E. Coligan, A. M.

25   Kruisbeek, D. H. Margulies, E. M. Shevach, and W. Strober, eds., Greene Publishing Associates, Brooklyn, NY. Cells were incubated with 1 uM ODN at 37°C for 8-12 h, and their secretion of cytokine detected colorimetrically as previously described. Klinman, D.M., and Nutman, T.B., *Current Protocols in Immunology* (1994), J. E. Coligan, A. M. Kruisbeek, D. H. Margulies, E. M.

30   Shevach, and W. Strober, eds., Greene Publishing Associates, Brooklyn, NY.

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Cell-surface binding and internalization of ODN:

　　　　Spleen cells (2 x 106/ml) were incubated with 1 uM of unlabeled and/or fluorescent- labeled ODN for 10' at 4°C (binding experiments) or at 37°C for 1 h (uptake experiments Gursel, M., *et al., J.Leuko.Biol.* (2001) in press. Cells were

5　　washed, fixed, and analyzed by FACScan (Becton Dickinson, San Jose, CA).

Statistical analysis:

　　　　Statistically significant differences between two groups were determined using the Wilcoxon Rank Sum Test. When comparing more than two groups,

10　　differences were determined using a 2-tailed non-parametric ANOVA with Dunn's post-test analysis. p values <0.05 were considered significant.

B. 　· 　*Mammalian DNA suppresses CpG DNA-induced immune activation*

　　　　Single stranded bacterial DNA and synthetic ODN containing unmethylated

15　　CpG motifs stimulate immune cells to mature, proliferate and produce cytokines, chemokines, and Ig. Klinman, D.M., *et al., Proc. Natl. Acad. Sci. USA* (1996) 93:2879; Roman, M., *et al., Nature Medicine* (1997) 3:849. (Roman *et al.,* (1997) *Nature Medicine* 3:849; Yamamoto *et al.,* (1992) *J. Immunol.* 148:4072; Krieg *et al.,* (1995) *Nature* 374:546). These effects can be blocked by "poly-G" and/or "GC"

20　　rich DNA motifs (Krieg *et al.,* (1998) *Proc.Natl.Acad.Sci.* 95:12631; Pisetsky *et al.,* (1995) *NY Acad. Sci.* 772:152). Scores of ODNs were synthesized and tested, and eventually several were identified that selectively inhibited CpG-induced immune responses. The two most active of these suppressive ODN (ODN1502 and ODNH154) were selected for detailed study. As seen in Fig. 4, suppressive ODN

25　　blocked a majority of the IFNγ production induced by bacterial DNA or CpG ODN (p < .01). Suppressive ODN were neither toxic nor broadly immunosuppressive, as they did not interfere with the mitogenic activity of LPS or Con A (Fig. 4).

　　　　The activity of suppressive ODNs was concentration dependent, with 50% suppression being achieved at a suppressive:CpG ODN ratio of approximately 1:3

30　　(Fig. 5). To examine the kinetics of this inhibition, suppressive ODN were added to BALB/c spleen cells at various times after CpG-induced stimulation. Maximal inhibition was observed when suppressive ODN were co-administered with CpG

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ODN, although statistically significant inhibition persisted when suppressive ODN were added up to 3 h later (Fig. 6). These findings suggest that CpG induced immune activation is an ongoing process, and can be inhibited after the stimulatory signal is delivered.

5   To test this conclusion, spleen cells were incubated with CpG ODN for various periods and cytokine production analyzed after 24 h. Cells stimulated with CpG DNA for 8 h produced 90% as much cytokine as cells stimulated continuously for 24 h (Fig. 7). Cells treated with CpG ODN for only 4 h produced half as much cytokine, while cells treated with CpG DNA for < 2 h showed only minimal

10   activation (Fig. 7). These findings support the conclusion that CpG-induced cellular activation is reversible for several hours.

C.   *Suppressive ODNs do not block CpG ODN uptake or induce the production of inhibitory factors*

15   The results described above indicate that CpG-induced immune activation can be reversed either by adding suppressive ODNs or by removing stimulatory ODNs. This suggests that suppressive ODNs block the ongoing uptake of CpG DNA. Yet FACS analysis demonstrated that neither cell surface binding nor internalization of FITC-labeled CpG ODN was significantly reduced by suppressive

20   ODN at concentrations that blocked cytokine production by approximately 75% (Fig. 8).

The possibility that suppressive motifs might induce the production of a factor that blocks CpG-dependent immune activation was then investigated. Initial studies established that BALB/c spleen cells pre-incubated with suppressive ODN

25   remained unresponsive to CpG-induced stimulation for several hours (Table I, line 3). If this non- responsive state was mediated by a soluble factor (or inhibitory cell-cell interactions) then cells pre-treated with suppressive ODN should block CpG-induced stimulation of naive spleen. As seen in Table I, cells treated with suppressive ODN had no significant effect on CpG dependent cytokine production

30   by fresh splenocytes.

43

Table I

Effect of mixing cells treated with suppressive versus stimulatory ODN

Cells pre-treated with suppressive ODN	Fresh cells	ODN added during culture	% maximal cytokine production	
			IL-6	IL-12
-	+	CpG	100 ± 13	100 ± 6
-	+	Control	3 ± 2	7 ± 2
+	-	CpG	9 ± 6	6 ± 2
+	-	Control	0 ± 0	0 ± 0
+	+	CpG	86 ± 16	105 ± 12
+	+	Control	0 ± 0	0 ± 0

5

BALB/c spleen cells were treated with 1 uM suppressive ODN_{1502} for 2 h and then washed (first column). These cells were added to naive splenocytes (second column) plus 1 uM of control ODN_{1471} or CpG ODN_{1555}. IL-6 and IL-12 levels in culture supernatants were measured by ELISA after 18 h. Results represent the average + SD of triplicate assays, each standardized to the response induced by bacterial DNA (62 pg/ml IL-6; 134 pg/ml

10   IL-12).

D.   Cellular recognition of suppressive versus stimulatory motifs

The above studies establish that suppressive motifs on one strand of DNA block the immune activation induced by stimulatory motifs on a different strand

15   (*i.e.*, trans suppression). To better understand the interaction between suppressive and stimulatory motifs, ODNs containing both were synthesized. In the simplest case, a 20-mer was constructed in which a CpG motif was placed immediately 5' to a suppressive motif (referred to as [CpG - Sup] ODN).

Experiments showed that this ODN was stimulatory, triggering murine

20   spleen cells to produce IL-6, IL-12 and IFNγ to the same extent as an ODN of the same length in which the suppressive motif was replaced by a 'control' sequence (*i.e.*, one that was neither stimulatory nor suppressive, Table II). [CpG - Sup] ODNs also failed to block the immune activation induced by an independent CpG ODN (Table II). These results suggest that a suppressive motif is inactive when located

25   immediately 3' to a CpG motif on the same strand of DNA. Similar results were obtained in studies of additional [CpG - Sup] ODNs that utilized different stimulatory and suppressive motifs.

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Table II

Effect of motif position on immunostimulatory activity

Location of motifs (5' → 3')	# of cytokine secreting cells		
	IL-6	IL-12	IFNg
CpG ODN *	79 ± 13	1980 ± 230	260 ± 40
[CpG – Sup] ODN*	72 ± 14	2080 ± 480	230 ± 60
[Sup-CpG] ODN	0±0	140±30	0±0
[CpG-Cont] ODN*	64±12	2210±130	284±34
[Cont-CpG] ODN*	80±11	1942±88	238±28
[Cont-Sup] ODN	8±2	184±34	36±8
[CpG-Sup] ODN+ Sup ODN	4±2	226±38	28±6
[Sup-CpG] ODN + CpG ODN*	7±3	250±32	34±9

5

10^6 BALB/c spleen cells were co-incubated with 1 uM of each ODN. Complex ODN (20 bp in length) were constructed from 10-mers containing suppressive (Sup), stimulatory (CpG) or control (Cont) motifs. The number of cytokine secreting cells/10^6 was determined by ELIspot after 24 h of stimulation. Equivalent results were derived using combinations of motifs derived from two different stimulatory (ODN_{1555} and ODN_{1466}),

10   control (ODN_{1471} and ODN_{1612}) and suppressive (ODN_{1502} and ODN_{H154}) ODN. Results represent the average ± SD of triplicate assays involving at least two ODN of each type.

* Stimulatory ODN, $p < .05$.

To better understand this phenomenon, longer ODNs were synthesized in

15   which the CpG and suppressive motifs were separated by progressively longer CT spacers. Adding a 5 base spacer generated an ODN that was still stimulatory (Table III). However, separating the motifs by >10 bases yielded ODNs that were suppressive, since they blocked the stimulatory activity of co-administered CpG ODNs (Table III).

45

Table III

Effect of distance between motifs on ODN activity

ODN	Cytokine producing cells (% maximum)		
	IL-6	IL-12	IFNg
CpG ODN *	100 ± 11	100 ± 7	100 ± 10
CpG ODN * + Cont ODN	97 ± 14	98 ± 9	100 ± 17
CpG ODN * + Sup ODN	16 ± 6	21 ± 6	18 ± 5
[CpG – Sup] ODN *	87 ± 12	$>100 \pm 14$	92 ± 14
[CpG – 5 bases – Sup] ODN *	$>100 \pm 4$	$>100 \pm 21$	$>100 \pm 22$
[CpG – 10 bases – Sup] ODN	38 ± 6	64 ± 15	42 ± 7
[CpG – 20 bases – Sup] ODN	7 ± 4	48 ± 13	24 ± 8
[CpG – 20 bases – Cont] ODN *	94 ± 7	$>100 \pm 14$	99 ± 11
[Sup – CpG] ODN	0 ± 0	0 ± 0	0 ± 0
[Sup – 20 bases – CpG] ODN	8 ± 5	9 ± 3	2 ± 1
[CpG – Sup] ODN * + CpG ODN *	$>100 \pm 16$	$>100 \pm 15$	98 ± 13
[CpG – 5 bases – Sup] ODN * + CpG ODN *	$>100 \pm 18$	$>100 + 11$	98 ± 20
[CpG – 10 bases – Sup] ODN + CpG ODN *	58 ± 7	75 ± 9	66 ± 9
[CpG – 20 bases – Sup] ODN + CpG ODN *	27 ± 5	26 ± 10	30 ± 8
[Sup – CpG] ODN + CpG ODN *	9 ± 4	11 ± 4	8 ± 5
[Sup – 20 CT – CpG] ODN + CpG ODN*	5 ± 1	9 ± 3	13 ± 2

BALB/c spleen cells were stimulated in vitro with 1 uM of each ODN, and the number of cells activated to secrete cytokine determined 8 h later by ELIspot. The percent of cells activated to secrete cytokine was calculated by the formula:

(# of cells activated by test ODN) - (background)/

(# of cells activated by CpG ODN) - (background) X 100%.

Two different control (ODN_{1471}, and ODN_{1612}), CpG (ODN_{1466} and ODN_{1555}) and suppressive (ODN_{1502} and ODN_{H154}) ODN gave similar results in these experiments.

Results represent the average of 2-4 assays/data point. Table II shows typical numbers of cytokine secreting cells/10^6.

* Stimulatory ODN, p <.05.

The trivial possibility that the CT spacer somehow reduced CpG activity was eliminated by substituting a "control" motif for the 3' suppressive motif. The resulting ODNs were fully stimulatory (Table III).

The impact of placing a suppressive motif 5' to a CpG motif was then examined. ODNs with a suppressive motif in the 5' position induced little or no immune activation, even when the CpG motif was shifted up to 20 bp downstream

46

from the suppressive motif (Tables II and III). This lack of activity could not be

attributed to the 3' location of the CpG motif, since CpG ODNs with a 'control'

sequence at the 5' end were immunostimulatory. All ODNs containing a suppressive

motif in the 5' position also inhibited co-administered CpG ODN (Tables II and III).

5   These findings suggest that the relative position of stimulatory and suppressive

motifs determines the immunomodulatory properties of DNA.

Example 3

Suppressive ODN permit higher levels of Transgene Expression

10   This example demonstrates that suppressive ODNs permit higher levels of

transgene to be expressed when the transgene vector contains immunostimulatory

CpG motifs.

Balb/C mice were injected intramuscularly with 30µg vector encoding

erythropoietin alone or together with 50 µg of each suppressive ODN. Suppressive

15   ODN was either co-injected at the time of vector injection or was given 3 days prior

to vector injection. Transgene expression was studied by measuring the hematocrit

levels from blood at various intervals. Data represent mean + S.D. for three animals.

Suppressive ODN permitted higher levels of transgene expression (Fig. 9).

Without being bound by theory, it is believed that this increase in transgene

20   expression is limited by immune activation triggered by immunostimulatory CpG

motifs in the erythropoietin construct. Administration of suppressive ODN inhibits

this immune activation and permits the transgene to be expressed at higher levels.

Example 4

Suppressive ODN blocks the activity of CpG motifs *in vivo*

25   This example shows that suppressive ODNs work in vivo to directly

counteract the activity of immunostimulatory CpG motifs.

The effect of Suppressive ODN on CpG ODN-induced immune protection

was studied. Balb/c mice were injected IP with 50µg of each ODN. Five days later,

30   animals were challenged IP with 10^3 LD50 of *Listeria monocytogenes*. Survival was

monitored for >3 weeks, although all animals that succumbed to infection died

47

within one week of challenge. Suppressive ODN blocked the activity of CpG motifs *in vivo* (Fig. 10).

Example 5

5   **Reduction of CpG-Induced Arthritis by Suppressive Oligodeoxynucleotides**

This example demonstrates the influence of CpG DNA and suppressive ODNs on the propensity of the host to develop arthritis, and indicates that the mechanism of suppressive ODN action is mediated by Cd11c positive cells.

10   *A.   General methods*

Animals:

Female BALB/c mice were obtained from the Jackson Laboratories (Bar Harbor, Maine). ACUC approved studies were conducted using 8-20 week old mice.

15

Oligodeoxynucleotides:

ODNs were synthesized at the CBER Core Facility. Sequences of the phosphorothioate ODN used were:

CpG: GCTAGACGTTAGCGT (SEQ ID NO: 30)

20   suppressive: CCTCAAGCTTGAGGGG (SEQ ID NO: 1)

control: GCTAGATGTTAGCGT. (SEQ ID NO: 31)

All ODN were free of detectable protein or endotoxin contamination.

Experimental protocol:

25   25 g of ODN in 6 μl of PBS was injected into the knee joint using a 30 gauge needle. In some studies, knees were re-injected with PBS or 25 μg of suppressive ODN 24-48 h after initial CpG ODN administration. Joint swelling was measured in the coronal plane using a micrometer caliper. Histologic analysis was performed by a blinded investigator on fixed, decalcified and paraffin embedded sections stained

30   with hematoxylin/eosin.

48

TNFα assays:

Single spleen cell suspensions were prepared in RPMI-1640 supplemented
with 5% heat inactivated fetal calf serum, 1.5 mM L-glutamine and 100 U/ml of
penicillin/streptomycin. 5 x 10^5 cells/well were cultured in flat-bottomed microtiter
5   plates (Costar, Corning, New York) with ODN for 72 hr. TNFα levels in culture
supernatants were measured by ELISA. In brief, 96-well Immulon H2B plates were
coated with anti-TNFα Ab (Genzyme, Cambridge, Massachusetts). Plates were
blocked with PBS - 1% BSA and overlaid with culture supernatants. Bound
cytokine was detected by the addition of biotin labeled anti-TNFα Ab (Genzyme,
10   Cambridge, Massachusetts) followed by phosphatase-conjugate avidin.

RT-PCR:

Total RNA was extracted from homogenized knees using TRIzol reagent
(GibCO Life Technologies, Gaithersburg, Maryland) as recommended by the
15   manufacturer. 5 ug of total RNA was reverse transcribed into cDNA, which was
assayed for TNFα (sense ATGAGCACAGAAAGCATGATC, antisense
TACAGGCTTGTCACTCGAATT, 275 bp) and β-actin (sense
GACATGGAGGAGTCTGGCACCACA, antisense
ATCTCCTGCTCGAAGTCTAGAGCAA , 440 bp) by PCR as previously described
20   (Takeshita *et al., Neuroreport* (2001) 12:3029-3032). Relative band intensity was
determined by ethidium bromide staining of 1% agarose gels using NIH-Image
software.

Statistical analysis:

25   Statistically significant differences between two groups were determined
using the Wilcoxon Rank Sum Test. When comparing more than two groups, a two-
tailed non-parametric ANOVA with Dunn's post-test analysis was used. Differences
in joint diameters were analyzed by repeated-measures ANOVA using the Proc
Mixed procedure from the Statistical Analysis System (SAS). p values <0.05 were
30   considered significant.

49

B.   Induction of arthritis by CpG ODN

Microbial infection of the gastrointestinal (GI) or genitourinary (GU) tracts
is associated with the development of reactive arthritis in humans. Evidence
suggests that bacterial DNA contributes to this process, since 1) bacterial DNA can

5   be detected in arthritic joints (Braun *et al., J. Rheumatol.* (2000) 27:2185-2192), and
2) bacterial DNA induces joint inflammation when injected into the knees of normal
mice (Deng *et al., Nat. Med.* (1999) 5:702-705). Deng *et al.* established that
immunostimulatory CpG motifs were the cause of this inflammation by showing
that CpG-containing oligodeoxynucleotides (ODN) induced disease in a manner

10   similar to that induced by purified bacterial DNA (Deng *et al., supra*). This effect is
consistent with the proinflammatory properties of CpG ODN, including their ability
to stimulate immune cells to proliferate, differentiate, and secrete proinflammatory
chemokines and cytokines (Deng *et al., supra*; Klinman *et al., Springer Semin.
Immunopathol.* (2000) 22:173-83).

15   Consistent with the findings of Deng *et al.* (Deng et al., *Arthritis Rheum.*
(2000) 43:356-364), BALB/c knees injected with CpG ODN developed
inflammatory arthritis within 24 hr that peaked after 3-7 days (Fig. 11). CpG-
induced arthritis was characterized by swelling (0.14 + 0.04 mm vs 0.02 + 0.02, p=
0.054) and histological changes that included perivascular infiltration by

20   mononuclear cells and hyperplasia of the synovial lining (Fig. 11). These
inflammatory effects were CpG-specific and localized, since no disease was
observed in contra-lateral knees injected with PBS or control ODN. Similar
swelling and histologic changes were observed in knees injected with bacterial
DNA.

25

C.   Suppressive ODN block the development of CpG-induced arthritis

To determine whether suppressive ODN prevent CpG mediated
inflammatory arthritis, knees were co-injected with 25 g of suppressive plus 25 g of
CpG ODN. The inclusion of suppressive ODN reduced swelling from 0.14 + 0.04

30   mm to 0.02 + 0.02 mm (p= 0.004) and the inflammatory score from 1.94 + 0.32 to
0.67 + 0.12 (p =0.018, Fig. 11). When joint inflammation was assessed by magnetic
resonance imaging, suppressive ODN reduced CpG ODN induced fluid

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accumulation from 95.4 + 8.2 MR-signal intensity units to 52.3 + 6.7 units (n=5,

p<0.001). These effects were specific, since co-administering PBS or control ODN

had no impact on CpG-induced arthritis (Fig. 11). In parallel studies, suppressive

ODN prevented the arthritis induced by bacterial DNA but not LPS (data not

5   shown).

D.   *Kinetics of the ODN anti-inflammatory effect*

To examine the kinetics of this anti-inflammatory effect, suppressive ODN

were administered 0, 24 and 48 h after CpG ODN injection. To control for the

10   effect of multiple injections, the contralateral knee was injected with PBS, and the

difference in swelling between the two joints evaluated daily. A significant

reduction in swelling was observed when joints were treated with suppressive ODN

up to two days after CpG administration (p=0.012, Fig. 11C). However, maximal

control of arthritis required early intervention (p =0.011, D0 vs D2).

15

E.   *Suppressive ODNs reduce intra-articular TNF production*

Previous studies established that the magnitude of CpG-induced arthritis

correlated with intra-articular TNFα levels (Deng *et al., Arthritis Res.* (2001) 3:48-

53). Consistent with TNFα playing a critical role in the disease process, TNFα KO

20   mice fail to develop CpG-induced arthritis (Deng *et al., Arthritis Rheum.* (2000)

43:356-364; Ronaghy *et al., J. Immunology* (2002) 168:51-56). To evaluate whether

suppressive ODN had an effect on TNFα production, BALB/c spleen cells were

stimulated *in vitro* with CpG + suppressive ODN. As seen in Fig. 12, suppressive

ODN reduced TNFα production in a dose-dependent manner, whereas control ODN

25   had no effect.

To monitor the *in vivo* effect of suppressive ODN on TNF production,

cytokine mRNA levels were measured in the joint. Consistent with earlier reports

(Deng *et al., Arthritis Rheum.* (2000) 43:356-364; Ronaghy *et al., J. Immunology*

(2002) 168:51-56; Kyburz *et al., Arthritis Rheum.* (2001) 44 (Suppl): S396), CpG

30   ODN up-regulated local TNF mRNA levels (Fig. 12C, D). Co-administering

suppressive ODN reduced TNFα mRNA by >50% (p<0.003, Fig. 12).

51

Example 6

Systemic Effect of Stimulatory and Suppressive Oligodeoxynucleotides on the Induction of Inflammatory Arthritis

A.   General methods

5   *Animals:*

Female BALB/c mice were obtained from the Jackson Laboratories (Bar

Harbor, Maine). The mice were used at 8-20 weeks of age and were housed under

specific pathogen free conditions. All experiments were approved by the CBER

Animal Use and Care Committee.

10

Oligonucleotides:

ODN used in this study had a phosphorothioate backbone, and were

synthesized in the CBER Core Facility. They contained <0.1 EU of endotoxin per

mg of ODN, as assessed by a Limulus amebocyte lysate assay (QCL-1000,

15   BioWhittaker). The sequence of the CpG ODN was GCTAGACGTTAGCGT (SEQ.

ID NO: 30), of the suppressive ODN: CCTCAAGCTTGAGGGG (SEQ. ID NO: 1),

and of the control ODN: GCTAGATGTTAGCGT (SEQ. ID NO: 31).

Experimental protocols:

20   Arthritis was induced as previously described in Zeuner, RA, *et al., Arthritis*

Rheum. (2002), in press. Briefly 1 or 25 μg of ODN in 6 μl of PBS was injected into

the knee joint using a 30 gauge needle. Joint swelling was measured in the coronal

plane using a micrometer caliper (General Tools Mfg CO, NY, NY). Mice were

euthanized on day 4 and the knees fixed, decalcified, sectioned, and stained with

25   hematoxylin/eosin prior to histologic examination. Scoring was performed by a

blinded investigator using a scale of 0-4. 0=absence of inflammation, 1=sparse,

localized perivascular infiltrate, 2=moderate infiltrate, 3=moderate-dense infiltrate

with synovial hyperplasia, 4=dense infiltrate with pronounced synovial hyperplasia.

In some experiments, donor mice were injected IP with 300 μg of ODN in

30   PBS. A single cell suspension prepared from the spleens of these mice was

prepared, and 20 x 106 transferred IV to naive littermate recipients. These spleen

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cells were enriched or depleted of various subpopulations using magnetic bead separation (Vario-Macs System, Miltenyi) as recommended by the manufacturer.

Statistical Analysis:

5   Differences between two groups determined using the Wilcoxon Rank Sum Test. Differences between multiple groups were evaluated using a two-tailed ANOVA on Ranks with Dunn's post test analysis. Differences in joint diameters were compared by repeated-measures ANOVA. P values <0.05 were considered significant. Data are presented as mean + SEM.F.

10

B.   *Effect of systemic CpG ODN on sensitivity of joints to inflammatory stimuli*

Previous studies established that intra-articular injection of bacterial DNA or CpG ODN induces arthritis in mice, characterized by an inflammatory cell infiltrate, perivascular accumulation of mononuclear cells, and hyperplasia of the synovial

15   lining Deng, GM, and Tarkowski A., *Arthritis Rheum*. (2000) 43:356-364. Since reactive arthritis in humans is associated with infection of the GI or GU tract rather than the joint, we explored whether CpG DNA in the systemic circulation might alter the sensitivity of joints to inflammatory stimuli.

To examine this possibility, normal BALB/c mice were injected IP with 300

20   ug of CpG DNA, and then challenged with a sub-arthritogenic dose of local CpG ODN. As seen in Fig. 14, normal mice and mice treated with PBS or control ODN did not develop arthritis (no inflammation, no swelling) when injected with 1 μg of CpG. In contrast, animals pre-treated with systemic CpG DNA developed significant joint swelling, a mononuclear cell infiltrate, and synovial hyperplasia

25   when challenged with the same dose of CpG ODN. These changes were triggered by exposure to local CpG DNA, as no inflammation developed when the joints of systemically treated animals were injected with PBS.

These findings indicate that CpG DNA in the systemic circulation (perhaps released by dying bacteria in the GI or GU tract) can increase the host's

30   susceptibility to the development local arthritis.

53

C.   *Systemic administration of suppressive ODN reduces susceptibility to arthritis*

It has been demonstrated that suppressive ODN (containing motifs that selectively inhibit the immunostimulatory activity of CpG ODN) significantly

5   reduce the swelling, synovial hyperplasia and leukocyte infiltration induced by CpG DNA. These effects were observed when suppressive ODN were injected directly into arthritic joints. Based on the observation that systemic CpG DNA can increase the host's susceptibility to arthritis, we postulated that suppressive ODN in the systemic circulation might lower this susceptibility.

10   To test this hypothesis, BALB/c mice were treated IP with 300 ug of suppressive ODN 0-3 days prior to the intra-articular injection of 25 ug CpG DNA. Consistent with previous studies, naive mice (and mice pre-treated with control ODN or PBS) developed severe arthritis following local CpG ODN challenge (Fig. 15). In contrast, systemic administration of suppressive ODN three days prior to

15   local CpG DNA challenge reduced joint swelling and inflammation by 80 - 85% (Fig. 16; $p<0.029$). Unlike the situation with locally administered suppressive ODN, which reduced inflammation when administered up to 2 days after the induction of arthritis, suppressive ODN were effective systemically only if delivered several days prior to the induction of arthritis. These findings suggest that instead of blocking

20   CpG induced arthritis locally, suppressive ODN in the systemic circulation might activate a regulatory cascade that requires several days to mature.

D.   *Spleen cells from suppressive ODN treated mice prevent CpG-induced arthritis*

25   To investigate this possibility, spleen cells from mice treated systemically with suppressive ODN were transferred to naive controls, which were then injected with 25 ug of CpG DNA intra-articularly. As expected, spleen cells from untreated donors had no effect on the development of CpG-induced arthritis (Fig. 17). By comparison, the transfer of 20×10^6 spleen cells from suppressive ODN treated

30   donors significantly reduced joint swelling and inflammation in the recipients. These findings indicate that systemically administered suppressive ODN elicit a population of regulatory cells that inhibit CpG-induced arthritis.

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To define the time course over which these regulatory cells are generated, splenocytes were isolated from 1-3 days after treatment with suppressive ODN. Cells from animals treated with suppressive ODN for 3 days significantly inhibited CpG-induced arthritis, whereas splenocytes from animals exposed to suppressive

5   ODN for shorter periods were progressively less effective.

To characterize the cell type responsible for this resistance to CpG-induced arthritis, magnetic beads were used to deplete or enrich specific cell subpopulations from donor spleens. Depletion of CD19+ B cells, T cells, or NK cells had no effect on CpG-induced arthritis (Fig. 18). However, removal of CD11c+ dendritic cells

10   resulted in a complete abrogation of the suppressive activity of the transferred spleen cell population. Similarly, the transfer of only 5 x 105 CD11c+ enriched cells from suppressive ODN treated mice to normal recipients conferred resistance to CpG-induced arthritis.

15   **Example 7**
                Neutralizing ODN Inhibits HSV DNA-Induced Angiogenesis
     A.   General Methods

Phosphorothioate ODNs were synthesized at the Center for Biologics Evaluation and Research Core Facility, as described (Verthelyi *et al.*, (2001) *J.*

20   *Immunol.* **166**, 2372-2377). The sequences of the stimulatory ODNs used in this study were: 1466, TCAACGTTGA (SEQ. ID NO: 26), and 1555, GCTAGACGTTAGCGT (SEQ ID No: 27); subsequent studies were conducted using an equimolar mixture of ODNs 1466 and 1555), the control ODN 1471 has the sequence TCAAGCTTGA (SEQ ID NO: 28), and the neutralizing ODN has the

25   sequence GAGCAAGCTGGACCTTCCAT (SEQ ID NO: 20). There was no detectable endotoxin contamination in any of the ODNs, as monitored by Limulus amebocyte lysate assay (BioWhittaker). In some experiments, FITC was conjugated to the 5' end of these ODN to monitor their distribution *in vivo*.

Herring sperm DNA (Boehringer Mannheim) was prepared by passage

30   through Detoxi-Gel Endotoxin Removal Gel 20344 (Pierce) to reduce endotoxin levels to <6 endotoxin units per mg. Recombinant human VEGF165 (rhVEGF), recombinant mouse VEGF (rmVEGF), mouse VEGF-neutralizing antibody, and biotinylated rat anti-mouse VEGF were purchased from R & D Systems. Synthetic

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mesh and hydron polymer (type NCC) were purchased from Sefar America (Kansas City, MO) and Hydro Med Sciences (Cranbury, NJ), respectively. Sucralfate was kindly provided by Bulch Meditec (Vaerlose, Denmark). Lipopolysaccharide was purchased from Sigma, and streptavidin-PE from PharMingen.

5

Isolation of HSV-1 DNA

Virus was harvested from infected Vero cells when the cytopathic effects were maximal, as described (Killington *et al.*, (1996) in Virology Methods Manual, eds. Mahy, B. W. J. & Kangro, H. O. (Academic, New York), pp. 71-88; McCance,

10   D. J. (1996) in Virology Methods Manual, eds. Mahy, B. W. J. & Kangro, H. O. (Academic, New York), pp. 191-197). Cells were suspended in sterile PBS and freeze-thawed three times to release viral particles. The virion-containing supernatant was then ultracentrifuged at 25,000 × g for 90 min at 4°C, and the pellet suspended in sterile PBS. Viral particles were precipitated in a solution of 7%

15   polyethylene glycol 8000 in 2.3% NaCl overnight at 4°C and then centrifuged at 25,000 × g. DNA was isolated from virions by treatment with 200 μg/ml proteinase K and 1% sarcosyl in STE buffer overnight at 56°C. The DNA was purified by multiple phenol/chloroform/isoamyl alcohol extractions, precipitated, sedimented at 12,000 × g, dried, and resuspended in sterile STE buffer. RNA was removed by

20   incubation with RNase (100 mg/ml; 5 Prime 3 Prime) for 1 h at 37°C, and the DNA reextracted as described above. The resultant material provided a single band by electrophoresis and contained undetectable protein (checked spectrophotometrically) or cellular DNA (measured by PCR for -actin DNA). All procedures were performed in a sterile environment, and all buffers and solutions were checked for

25   the presence of lipopolysaccharide by using the Pyrogent plus test. No detectable protein, cellular RNA, or DNA, and less than 0.06 endotoxin units of endotoxin per mg of HSV DNA, were found.

Mice

30   Female BALB/c (Harlan-Sprague-Dawley) were used for all experiments. Animals were housed and cared for as described (Gangappa *et al.*, (1998) *J. Immunol.* **161**, 4289-4300).

56

Corneal Micropocket Assay

The murine corneal micropocket assay used in this work observed the general protocol of Kenyon *et al.* (Kenyon *et al.,* (1996) *Invest. Ophthalmol. Vis.*

5   *Sci.* 37, 1625-1632). Pellets 0.4 × 0.4 × 0.2 mm3 composed of sucralfate and hydron polymer were prepared (Kenyon *et al.,* (1996) *Invest. Ophthalmol. Vis. Sci.* 37, 1625-1632). Known amounts of VEGF, DNA, stimulatory or neutralizing ODN, and/or combinations thereof were added to these pellets before insertion into corneal pockets. The micropockets were placed 0.6-0.8 mm from the limbus (in the

10   pericenter of the cornea at the lateral canthus of the eye) under stereomicroscopy (four eyes per group). In some experiments, anti-mVEGF-neutralizing antibody (5 μg in 5 μl of PBS) was injected subconjunctivally into the eyes of recipient mice just before and 2 days after pellet implantation.

Angiogenesis was quantitated at multiple times after pellet implantation

15   under stereomicroscopy. The length of the neovessels generated from the limbal vessel ring toward the center of the cornea and the width of the neovessels presented in clock hours (each clock hour is equal to 30°C at the circumference) was measured (Zheng *et al.,* (2001) *Am. J. Pathol.* 159, 1021-1029). The angiogenic area was calculated according to the formula for an ellipse. A = [(clock hours) × 0.4 × (vessel

20   length in mm) ×]/2.

Immunohistochemical Staining

Eyes were removed and snap frozen in OCT compound (Miles). Sections (6-μm) were cut, air dried, and fixed in cold acetone for 10 min. The sections were

25   blocked with 3% BSA and stained with biotinylated anti-mVEGF164. Sections were then treated with horseradish peroxidase-conjugated streptavidin (1:1,000) and 3,3'-diaminobenzidine (Vector), and counterstained with hematoxylin as described (Zheng *et al.,* (2001) *J. Virol.* 75, 9828-9835). Cellular infiltration was determined microscopically by counting the infiltrating cells in the corneal stroma. Each data

30   point represents the mean total cellular infiltrate in four central corneal sections from two eyes.

57

VEGF Staining of J774A.1 Cells

        J774A.1 cells were plated and incubated in two-well chamber slides (Lab-
Tek, Nalge Nunc International) or in 24-well plates [for later reverse transcription
(RT)-PCR] in DMEM with 10% FBS overnight at 37°C in 5% CO_2. The cells in
5   chamber slides were cocultured with FITC-labeled CpG ODN (1555) or control
ODN (1471) at a concentration of $2 \, \mu g/10^6$ cells. The cells were washed twice with
PBS and fixed in a 1:1 mixture of acetone/methyl alcohol at 20°C for 15 min. The
cells were stained with biotinylated rat-anti-mVEGF 6-18 h after ODN stimulation
and subsequently reacted with streptavidin-PE. Images were taken by using a
10 fluorescence microscope (Hamamatsu, Ichinocho, Japan). The cells in 24-well
plates were treated with $2 \, \mu g$ of ODN per 10^6 cells per ml. RNA from these cells
was extracted for RT-PCR to detect VEGF mRNA (see RNA Extraction and RT-
PCR).

15 *Fluorescence-Activated Cell Sorter Staining of VEGF-Expressing Cells*

        J774A.1 cells were treated with 0-8 $\mu g/ml$ of ODN for 6-12 h. The cells
were then fixed in paraformaldehyde, blocked with FCS, and stained for VEGF by
using biotinylated rat-anti-mVEGF164 antibody followed by streptavidin-PE.
Positive cells were identified by flow cytometry.

20

Statistical Analysis

        Significant differences between groups were evaluated by using the Student's
t test. P 0.05 was regarded as significant difference between two groups.

25 B.   *Neutralizing ODN Inhibits HSV DNA-Induced Angiogenesis*

        To investigate further whether CpG motifs in HSV DNA were responsible
for the angiogenesis, HSV DNA was mixed with "neutralizing" or control ODN and
the effect on angiogenesis recorded. The neutralizing ODN contained multiple G-
tetrad-forming motifs that other studies have shown to neutralize the
30 immunostimulatory effects of CpG DNA (Krieg *et al.,* (1998) Proc. Natl. Acad. Sci.
USA 95, 12631-12636; Lenert *et al.,* (2001) Antisense Nucleic Acid Drug Dev. 11,
247-256; Pisetsky *et al.,* (2000) Clin. Immunol. 96, 198-204). In these experiments,

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hydron pellets were prepared that contained HSV DNA along with neutralizing or control ODN. Levels of angiogenesis were recorded at 4 days after implantation. As is evident, pellets that incorporated control ODN plus HSV DNA induced the same level of angiogenesis as did HSV DNA pellets alone. In contrast, pellets

5   containing neutralizing ODN plus HSV DNA induced significantly less (60% inhibition) angiogenesis ($P < 0.01$, Fig. 19).

While this disclosure has been described with an emphasis upon preferred embodiments, it will be obvious to those of ordinary skill in the art that variations of

10   the preferred embodiments may be used and it is intended that the disclosure may be practiced otherwise than as specifically described herein. Accordingly, this disclosure includes all modifications encompassed within the spirit and scope of the disclosure as defined by the following claims:

59

We claim:

1. A substantially pure or isolated oligodeoxynucleotide of at least about 8 nucleotides in length, wherein:

    the oligodeoxynucleotide forms a G-tetrad;

5   the oligodeoxynucleotide has a CD value of greater than about 2.9;

    the number of guanosines is at least two; and

    wherein the oligodeoxynucleotide suppresses an immune response.

2. The oligodeoxynucleotide of claim 1, wherein the oligodeoxynucleotide

10 has a CD value of greater than about 3.0.

3. The oligodeoxynucleotide of claim 1, wherein the oligodeoxynucleotide has a CD value of greater than about 3.2.

15 4. The oligodeoxynucleotide of claim 1, wherein the oligodeoxynucleotide is from about 8 to about 120 nucleotides in length.

5. The oligodeoxynucleotide of claim 1, wherein the oligodeoxynucleotide is from about 10 to about 30 nucleotides in length.

20

6. The oligodeoxynucleotide of claim 1, wherein the oligodeoxynucleotide comprises multiple guanosine-rich sequences.

7. The oligodeoxynucleotide of claim 6, wherein the oligodeoxynucleotide

25 comprises from about two to about 20 guanosine-rich sequences.

8. The oligodeoxynucleotide of claim 6, wherein the oligodeoxynucleotide comprises from about two to about four guanosine-rich sequences.

30 9. The oligodeoxynucleotide of claim 1, wherein the oligodeoxynucleotide comprises at least one TTAGGG motif.

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10. The oligodeoxynucleotide of claim 9, wherein the oligodeoxynucleotide comprises multiple TTAGGG motifs.

11. The oligodeoxynucleotide of claim 9, wherein the oligodeoxynucleotide
5   comprises from about two to about 20 TTAGGG motifs.

12. The oligodeoxynucleotide of claim 9, wherein the oligodeoxynucleotide comprises from about two to about four TTAGGG motifs.

10   13. The oligodeoxynucleotide of claim 1, wherein the oligodeoxynucleotide comprises a sequence selected from the group consisting of SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 7, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO: 10, SEQ ID NO: 11, SEQ ID NO: 12, SEQ ID NO: 13, SEQ ID NO: 14, SEQ ID NO: 15, SEQ ID NO: 16, SEQ ID
15   NO: 17, SEQ ID NO: 18, SEQ ID NO: 19, SEQ ID NO: 20, SEQ ID NO: 21, SEQ ID NO: 22, SEQ ID NO: 23, SEQ ID NO: 24, and SEQ ID NO: 25.

14. The oligodeoxynucleotide of claim 1, wherein the oligodeoxynucleotide is modified to prevent degradation.

15. The oligodeoxynucleotide of claim 1, wherein the oligodeoxynucleotide
20   has a phosphate backbone modification.

16. The oligodeoxynucleotide of claim 15, wherein the phosphate backbone modification is a phosphorothioate backbone modification.

17. The oligodeoxynucleotide of claim 1, wherein the oligodeoxynucleotide suppresses an immune response induced by an immunostimulatory CpG
25   oligodeoxynucleotide.

18. An oligodeoxynucleotide delivery complex comprising the oligodeoxynucleotide of claim 1 and a targeting moiety.

19. The oligodeoxynucleotide delivery complex of claim 18, wherein the targeting moiety is selected from the group consisting of a cholesterol, a virosome, a liposome, a lipid, and a target cell specific binding agent.

20. The oligodeoxynucleotide of delivery complex of claim 18, wherein the oligodeoxynucleotide and the targeting moiety are covalently linked.

21. A pharmacological composition comprising the oligodeoxynucleotide of claim 1 and a pharmacologically acceptable carrier.

22. A method of suppressing an immune response in a subject comprising administering a therapeutically effective amount of the oligodeoxynucleotide of claim 1 to a subject in which it is desirable to suppress an immune response, thereby suppressing the immune response.

23. The method of claim 22, wherein the oligodeoxynucleotide is administered topically, parenterally, orally, intravenously, intra-muscularly, sub-cutaneously, inhalationally, nasally, topically, or intra-articularly.

24. The method of claim 22, wherein the oligodeoxynucleotide has a CD value of greater than about 3.0.

25. The method of claim 22, wherein the oligodeoxynucleotide has a CD value of greater than about 3.2.

26. The method of claim 22, wherein the oligodeoxynucleotide is from about 8 to about 120 nucleotides in length.

27. The method of claim 22, wherein the oligodeoxynucleotide is from about 10 to about 30 nucleotides in length.

28. The method of claim 22, wherein the oligodeoxynucleotide comprises multiple guanosine-rich sequences.

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29. The method of claim 22, wherein the oligodeoxynucleotide comprises from about two to about 20 guanosine-rich sequences.

5   30. The method of claim 22, wherein the oligodeoxynucleotide comprises from about two to about four guanosine-rich sequences.

31. The method of claim 22, wherein the oligodeoxynucleotide comprises at least one TTAGGG motif.

10

32. The method of claim 22, wherein the oligodeoxynucleotide comprises multiple TTAGGG motifs.

33. The method of claim 22, wherein the oligodeoxynucleotide comprises

15   from about two to about 20 TTAGGG motifs.

34. The method of claim 22, wherein the oligodeoxynucleotide comprises from about two to about four TTAGGG motifs.

20   35. The method of claim 22, wherein the oligodeoxynucleotide comprises a sequence selected from the group consisting of SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 7, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO: 10, SEQ ID NO: 11, SEQ ID NO: 12, SEQ ID NO: 13, SEQ ID NO: 14, SEQ ID NO: 15, SEQ ID NO: 16, SEQ ID NO: 17, SEQ

25   ID NO: 18, SEQ ID NO: 19, SEQ ID NO: 20, SEQ ID NO: 21, SEQ ID NO: 22, SEQ ID NO: 23, SEQ ID NO: 24, and SEQ ID NO: 25.

36. The method of claim 22, further comprising administering an additional anti-inflammatory or immunosuppressive agent.

30

37. The method of claim 36, wherein the agent is diclofenac, diflunisal, etodolac, flurbiprofen, ibuprofen, indomethacin, ketoprofen, ketorolac, nabumetone,

63

naproxen, oxaprozin, piroxicam, sulindac, tolmetin, celecoxib, rofecoxib, cortisone, dexamethasone, hydrocortisone, methylprednisolone, prednisolone, prednisone, triamcinolone, cyclosporin, tacrolimus, mycophenolic acid, or sirolimus.

38. A method of treating or preventing an immune-mediated disorder in a subject comprising contacting immune cells with the oligodeoxynucleotide of claim 1, and transferring the immune cells to a subject having or at risk of developing an immune-mediated disorder, thereby treating or preventing the immune-mediated disorder.

39. The method of claim 38, wherein the immune cells are transplanted.

40. The method of claim 38, wherein the immune cells are transferred intravenously.

41. A method of treating or preventing an immune-mediated disorder in a subject comprising administering a therapeutically effective amount of the oligodeoxynucleotide of claim 1 to a subject having or at risk for developing the immune-mediated disorder, thereby treating or preventing an immune-mediated disorder.

42. The method of claim 41, wherein the immune-mediated disorder is an autoimmune disease or graft-versus-host disease.

43. The method of claim 41, wherein the immune-mediated disorder is an autoimmune disease.

44. The method of claim 41, wherein the immune-mediated disorder is an inflammatory arthropathy.

45. The method of claim 44, wherein the oligodeoxynucleotide is administered topically, parenterally, orally, intravenously, intra-muscularly, sub-cutaneously, inhalationally, nasally, topically, or intra-articularly.

64

46. The method of claim 44, wherein the oligodeoxynucleotide has a CD value of greater than about 3.0.

5   47. The method of claim 44, wherein the oligodeoxynucleotide has a CD value of greater than about 3.2.

48. The method of claim 44, wherein the oligodeoxynucleotide is from about 8 to about 120 nucleotides in length.

10

49. The method of claim 44, wherein the oligodeoxynucleotide is from about 10 to about 30 nucleotides in length.

50. The method of claim 44, wherein the oligodeoxynucleotide comprises

15   multiple guanosine-rich sequences.

51. The method of claim 44, wherein the oligodeoxynucleotide comprises from about two to about 20 guanosine-rich sequences.

20   52. The method of claim 44, wherein the oligodeoxynucleotide comprises from about two to about four guanosine-rich sequences.

53. The method of claim 44, wherein the oligodeoxynucleotide comprises at least one TTAGGG motif.

25

54. The method of claim 44, wherein the oligodeoxynucleotide comprises multiple TTAGGG motifs.

55. The method of claim 44, wherein the oligodeoxynucleotide comprises

30   from about two to about 20 TTAGGG motifs.

65

56. The method of claim 44, wherein the oligodeoxynucleotide comprises from about two to about four TTAGGG motifs.

57. The method of claim 44, wherein the oligodeoxynucleotide comprises a
5   sequence selected from the group consisting of SEQ ID NO: 1, SEQ ID NO: 2, SEQ ID NO: 3, SEQ ID NO: 4, SEQ ID NO: 5, SEQ ID NO: 6, SEQ ID NO: 7, SEQ ID NO: 8, SEQ ID NO: 9, SEQ ID NO: 10, SEQ ID NO: 11, SEQ ID NO: 12, SEQ ID NO: 13, SEQ ID NO: 14, SEQ ID NO: 15, SEQ ID NO: 16, SEQ ID NO: 17, SEQ ID NO: 18, SEQ ID NO: 19, SEQ ID NO: 20, SEQ ID NO: 21, SEQ ID NO: 22,
10 SEQ ID NO: 23, SEQ ID NO: 24, and SEQ ID NO: 25.

58. The method of claim 44, further comprising administering an additional anti-inflammatory, immunosuppressive, or anti-arthritis agent.

15 59. The method of claim 58, wherein the agent is a biological response modifier, a disease-modifying antirheumatic drug (DMARD), a nonsteroidal anti-inflammatory drug (NSAIDs), or a Cyclo-Oxygenase-2 (COX-2) inhibitor.

60. The method of claim 58, wherein the agent is anakinra, etanercept, infliximab, leflunomide, celecoxib, rofecoxib, hyaluronan, or hylan G-F20.

20 61. A method of inhibiting angiogenesis in a subject comprising administering a therapeutically effective amount of the oligodeoxynucleotide of claim 1 to the, thereby inhibiting angiogenesis.

62. The method of claim 61, wherein the subject has a tumor.

63. The method of claim 61, wherein the administration is subconjunctival
25 administration.

64. A kit for treating or preventing autoimmune diseases in a subject comprising a container comprising the oligodeoxynucleotide of claim 1.

65. The kit of claim 64, further comprising a container comprising an additional anti-inflammatory or immunosuppressive agent.

66. The kit of claim 65, wherein the agent is diclofenac, diflunisal, etodolac, flurbiprofen, ibuprofen, indomethacin, ketoprofen, ketorolac, nabumetone,

5   naproxen, oxaprozin, piroxicam, sulindac, tolmetin, celecoxib, rofecoxib, cortisone, dexamethasone, hydrocortisone, methylprednisolone, prednisolone, prednisone, triamcinolone, cyclosporin, tacrolimus, mycophenolic acid, or sirolimus.

67. The kit of claim 64, further comprising instructions for administering the oligodeoxynucleotide to a subject
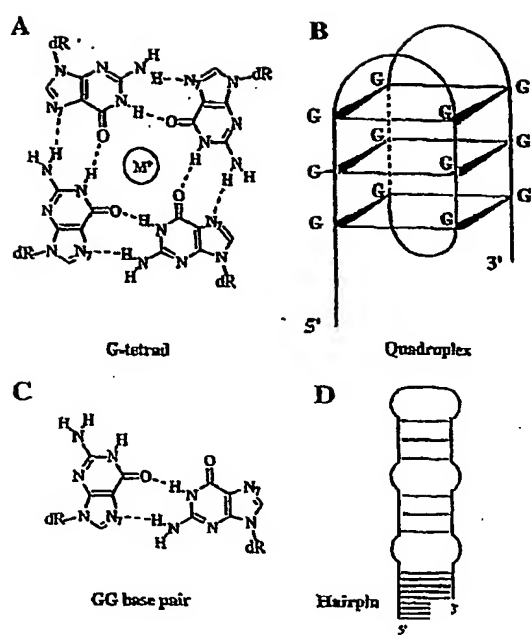
Fig. 1

a Mammalian DNA suppresses CpG DNA induced immune activation
  Bacterial DNA
  Mammalian DNA
  Bacterial DNA + Mammalian DNA
  CpG ODN
  CpG ODN + Bacterial DNA
  CpG ODN + Mammalian DNA
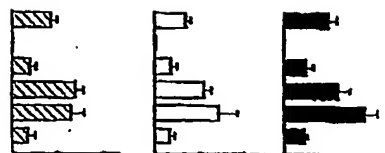
b Telomeric TTAGGG repetitive motif is suppressive

	Ellipticity (mdeg/abs)
CpG ODN	1.4
CpG ODN + (TTAGGG)₄	7.1
CpG ODN + (TTAGGG)₃	4.8
CpG ODN + (TTAGGG)₂	2.6
CpG ODN + (TTAGGG)₁	1.2
CpG ODN + TCAACCTTCATTAGGG	3.4
CpG ODN + TTAGGGTCAACCTTCA	3.4
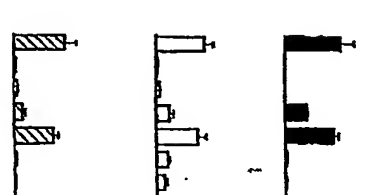
c Suppressive motif is active in *trans* and *cis*
  Trans
  CpG ODN
  CpG ODN + Control ODN
  CpG ODN + Suppressive ODN
  Cis
  (Control-CpG) ODN
  (Suppressive-CpG) ODN
  (Control-CpG) ODN + CpG ODN
  (Suppressive-CpG) ODN + CpG ODN
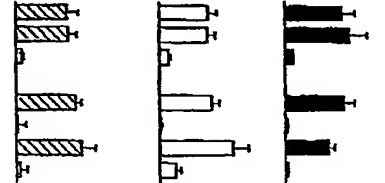
d Suppressive ODN selectively block CpG DNA induced immune activation
  CpG ODN
  CpG ODN + Suppressive ODN
  Bacterial DNA
  Bacterial DNA + Suppressive ODN
  LPS
  LPS + Suppressive ODN
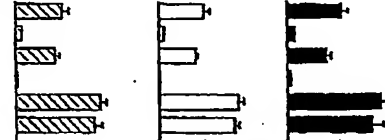
e Poly G's are critical for suppression

CpG ODN		1.4
CpG ODN +	TTAG G G TCAACCTTCA	3.4
CpG ODN +	– A C– – – – – – – – – – – –	
CpG ODN +	A C T – – – – – – – – – – – –	
CpG ODN +	A A T – – – – – – – – – – – –	
CpG ODN +	– – – – A A – – – – – – – – – –	
CpG ODN +	– – – T T – – – – – – – – – –	
CpG ODN +	– – – ᵀG– – – – – – – – –	1.2
CpG ODN +	– – – ᵀGᵀG– – – – – – – – –	1.0
CpG ODN +	– – – ᵀGᵀGᵀG– – – – – – – –	0.8
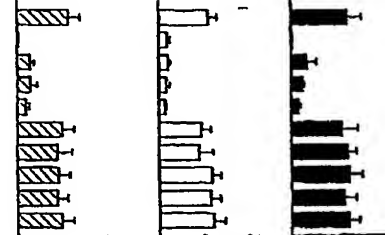
f G-tetrad forming non-telomere sequences are also suppressive

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CpG ODN +	CCT CAAGCTT GAGGGG	4.6
CpG ODN +	CCGGCCGGCCGGCCGG	2.8
CpG ODN +	TT GGTT GGTT GGTT GG	3.2
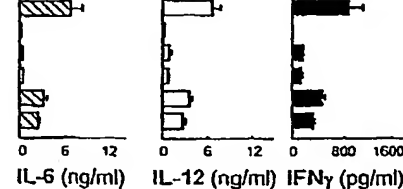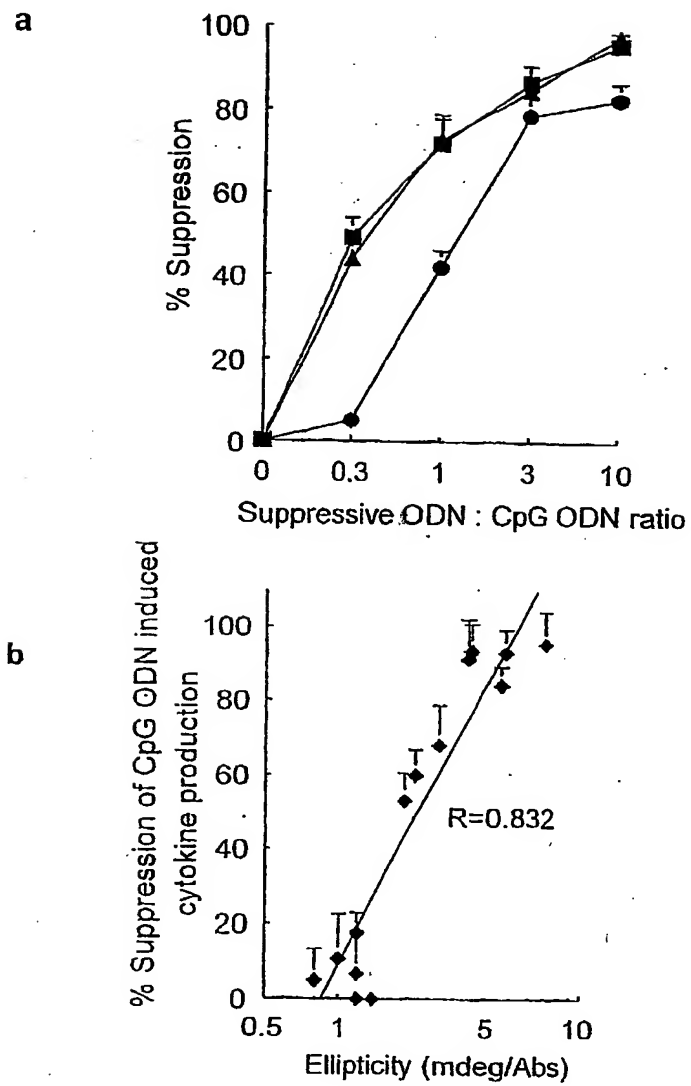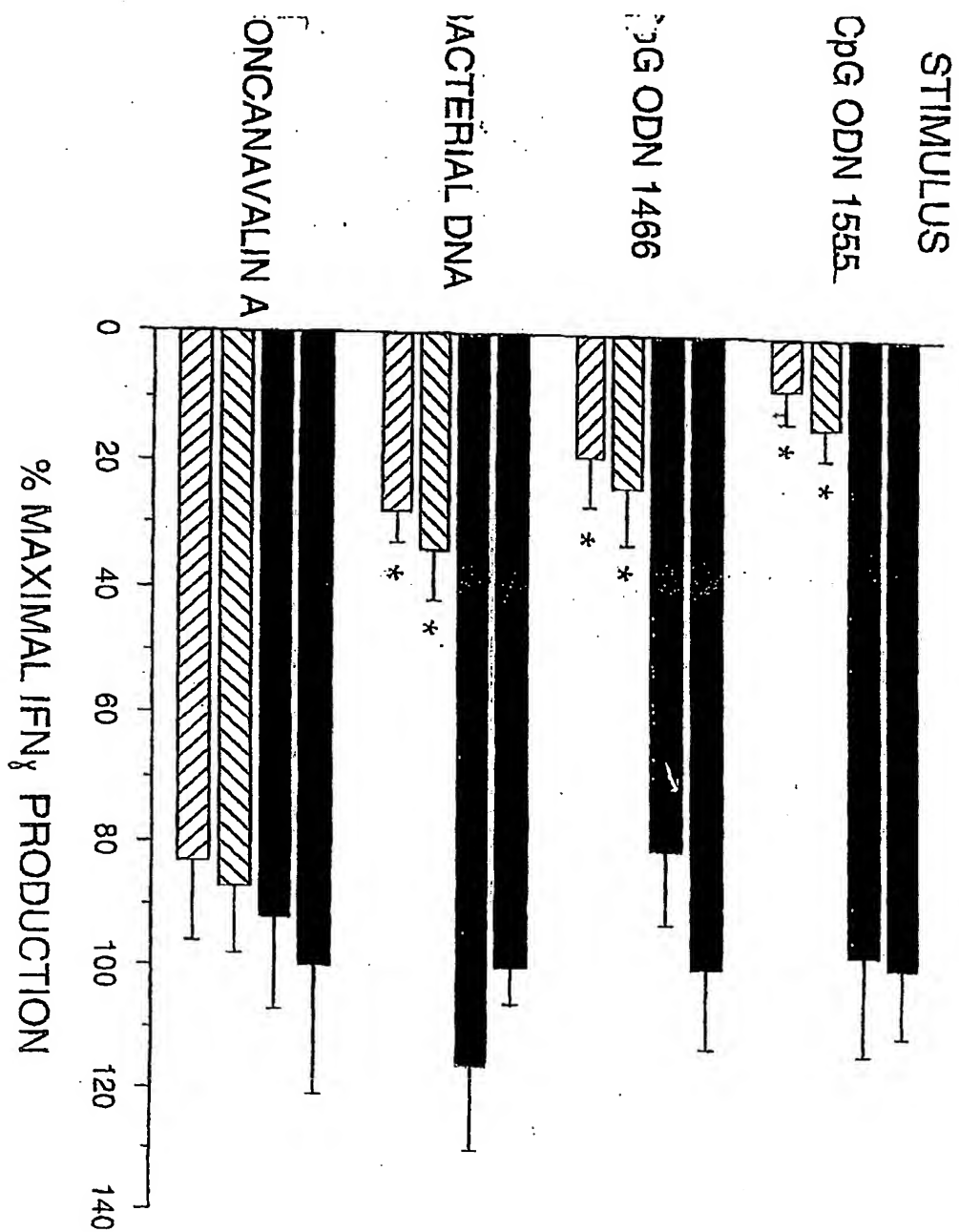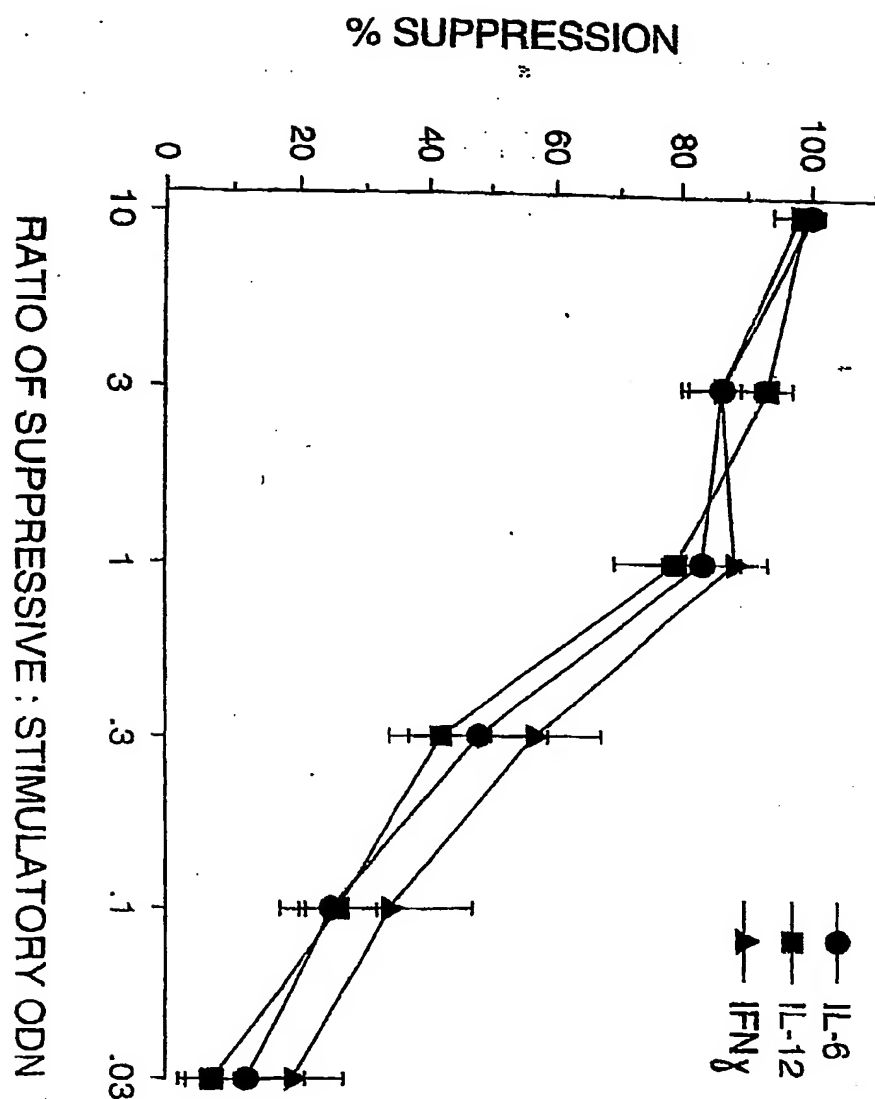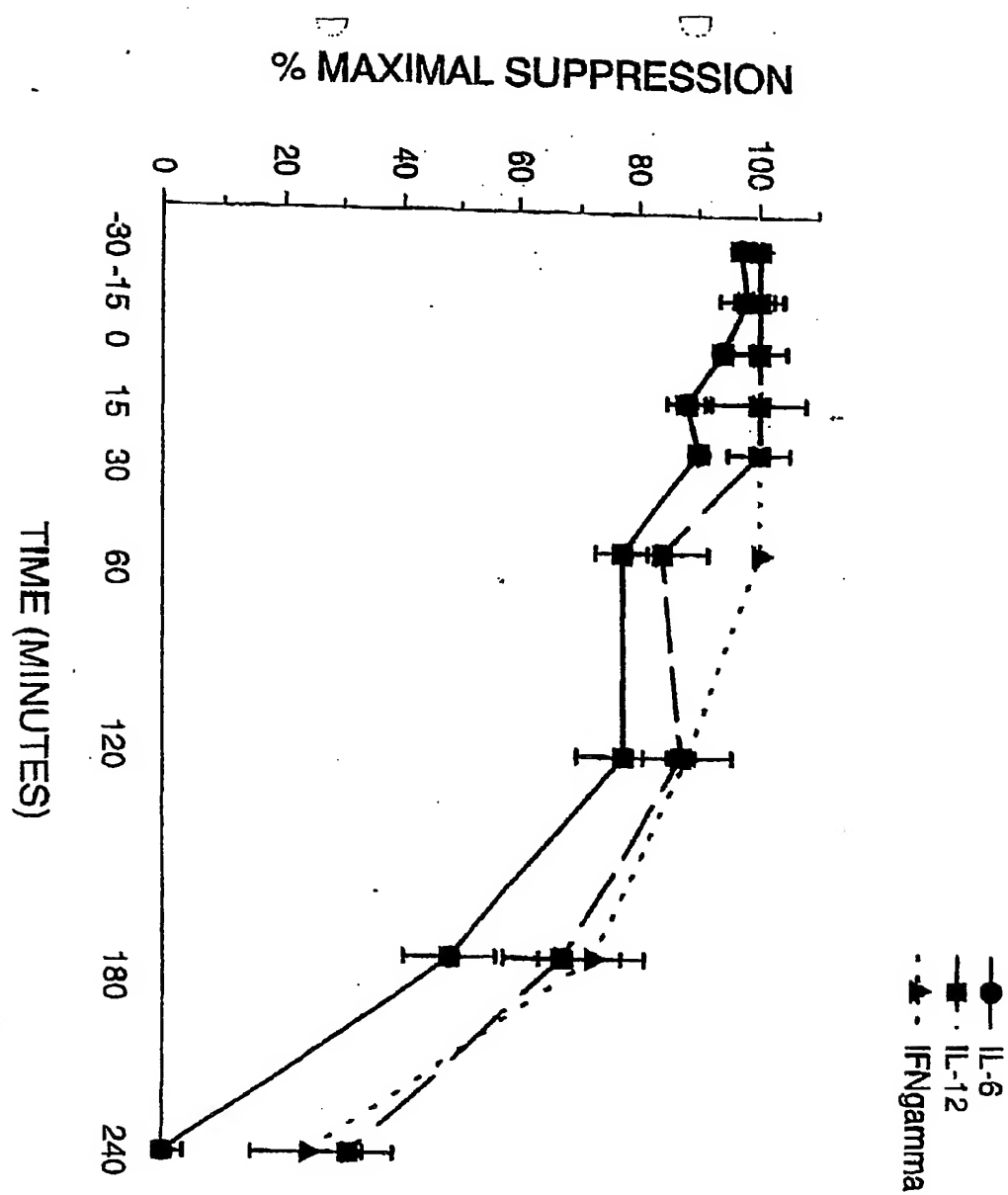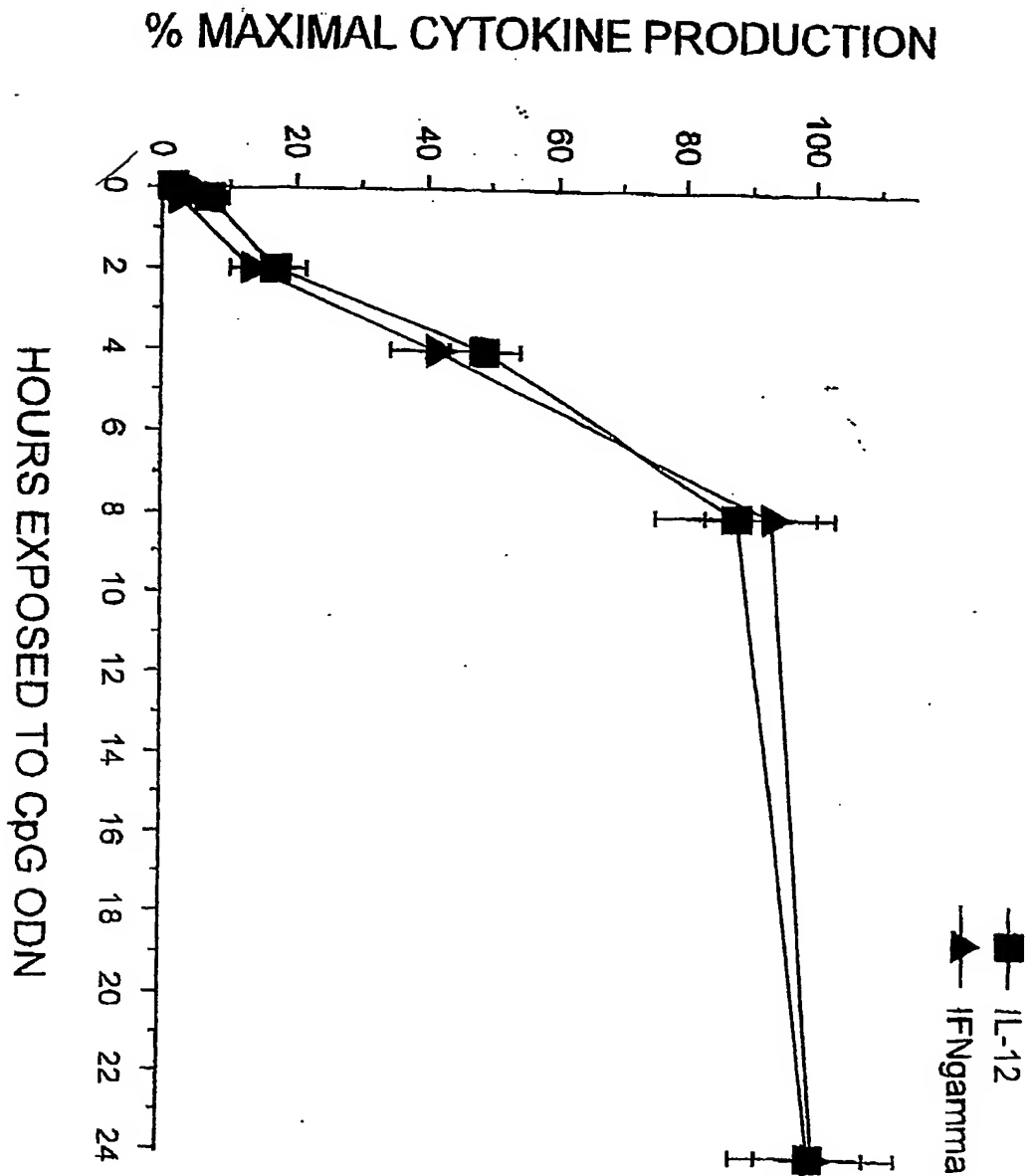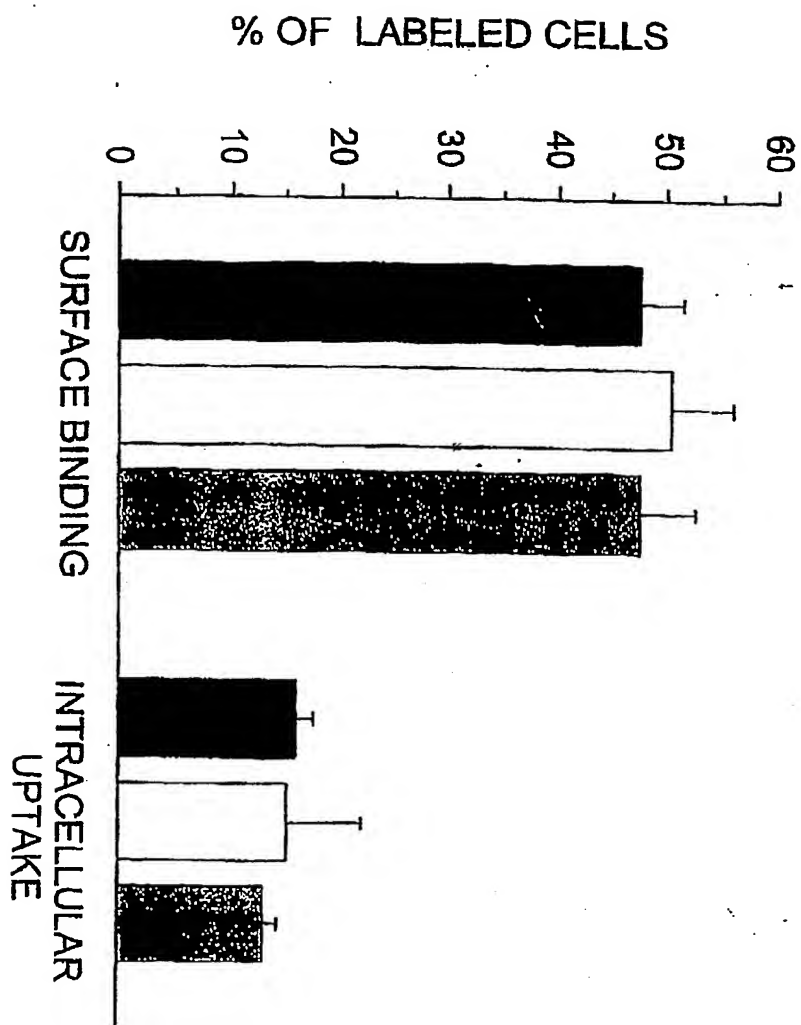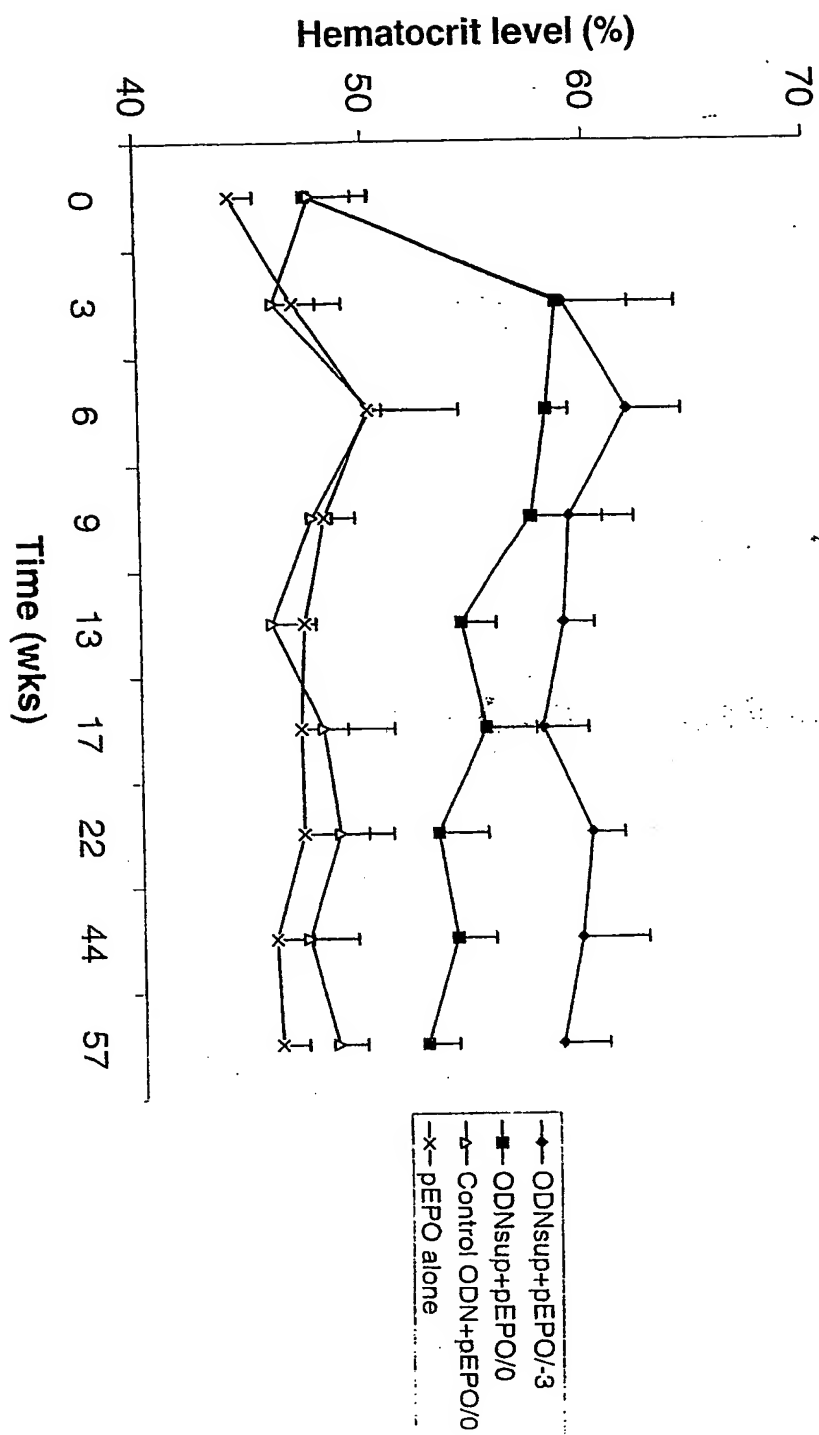
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IL-6 (ng/ml)   IL-12 (ng/ml) IFNγ (pg/ml)

Fig. 2

a

b

Fig. 3

Fig. 4

Fig. 5

Fig. 6

Fig. 7

Fig. 8

Fig. 9
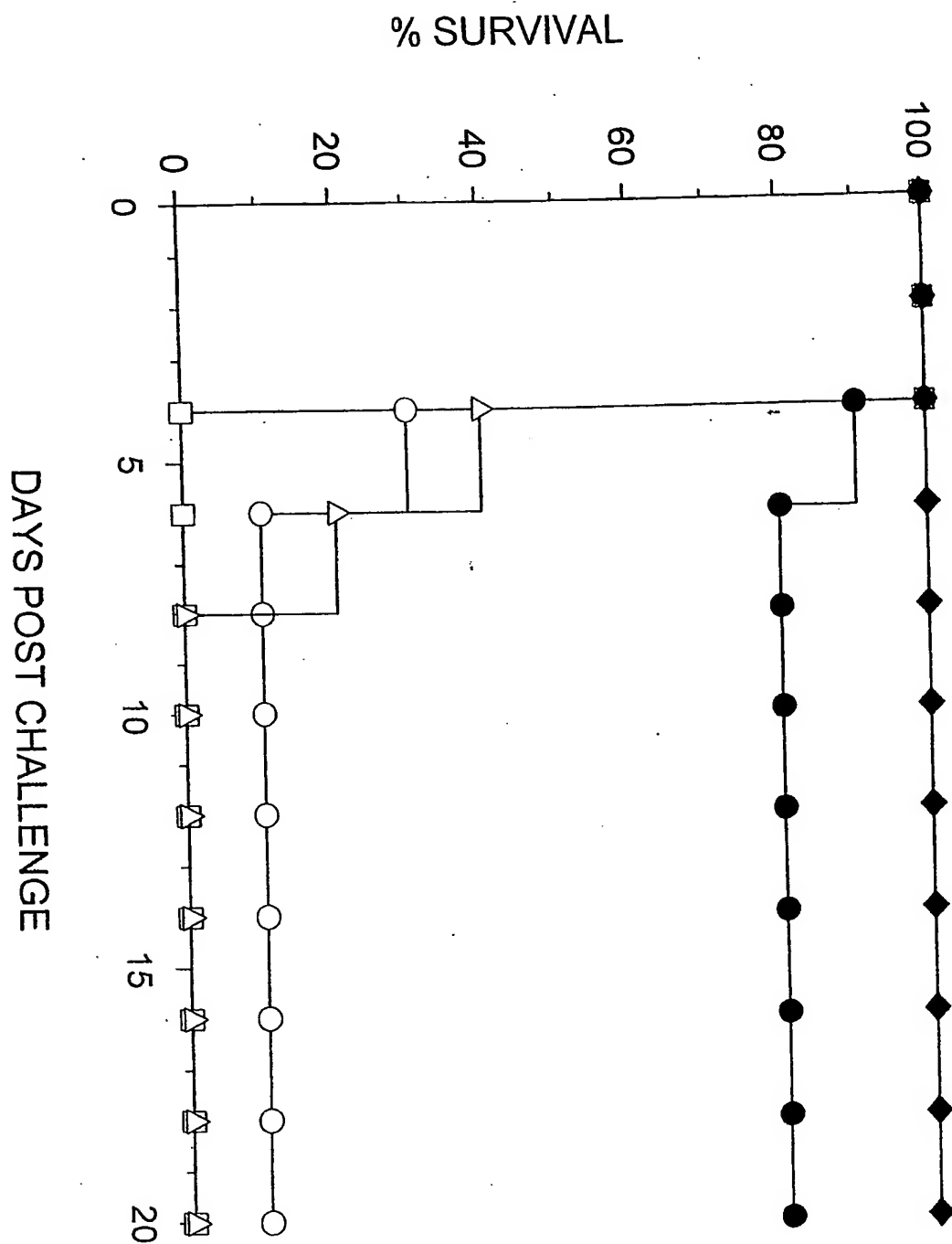
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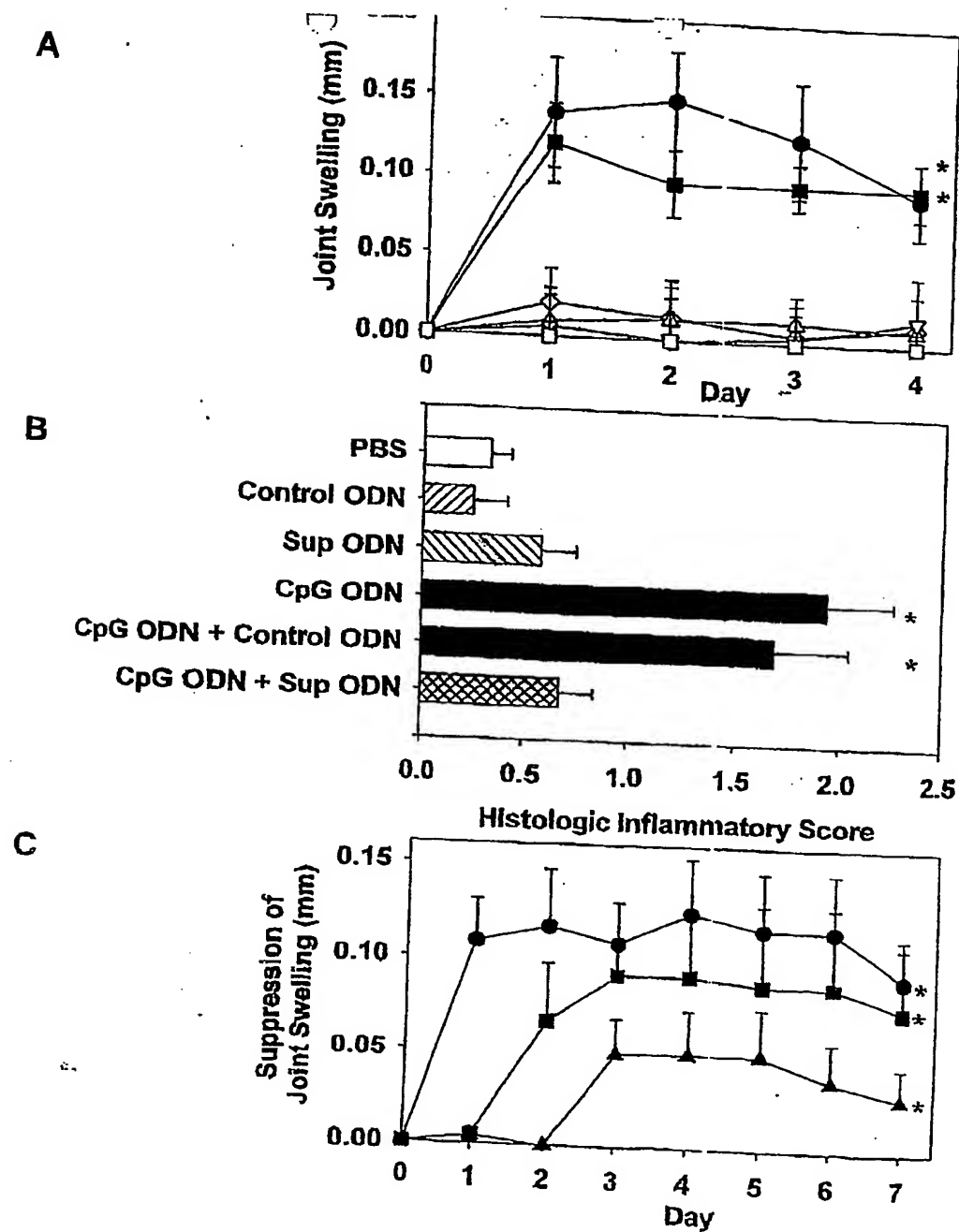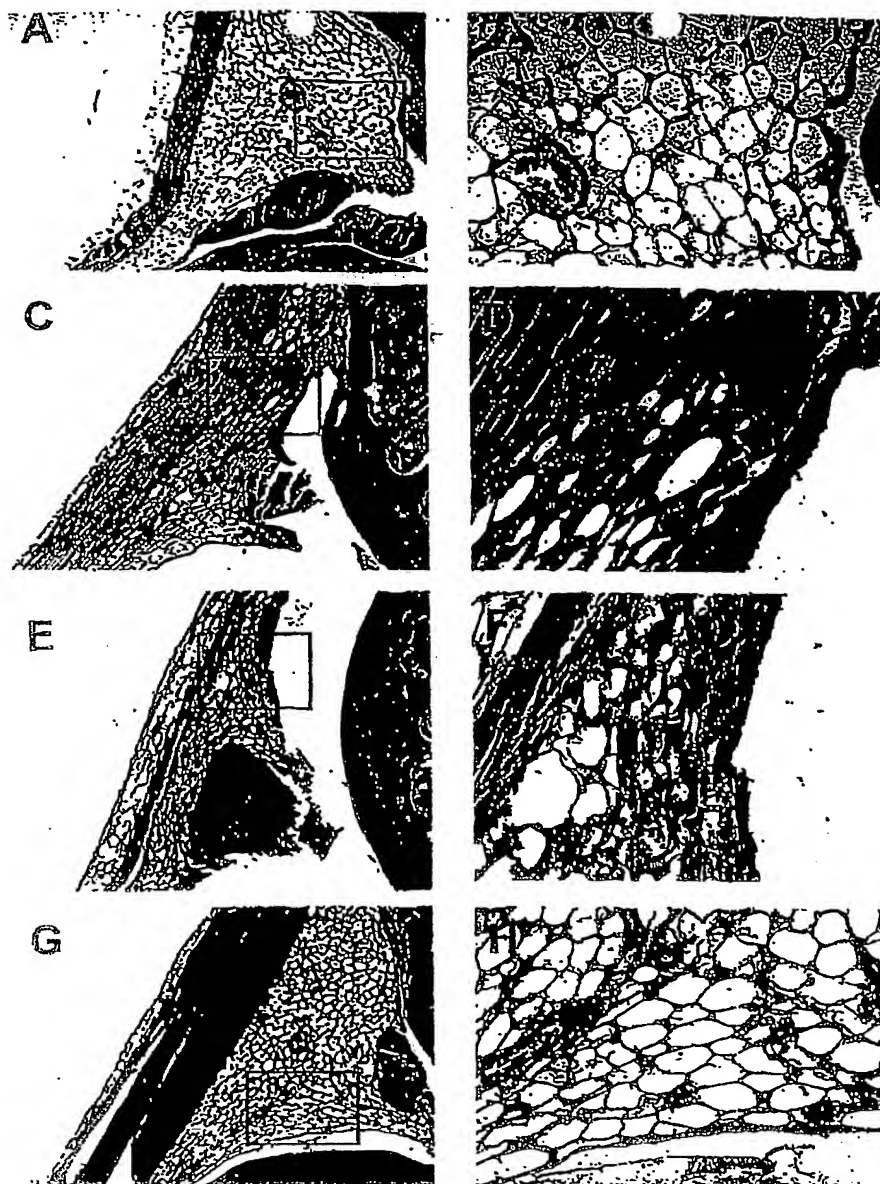
Fig. 11

Fig. 12
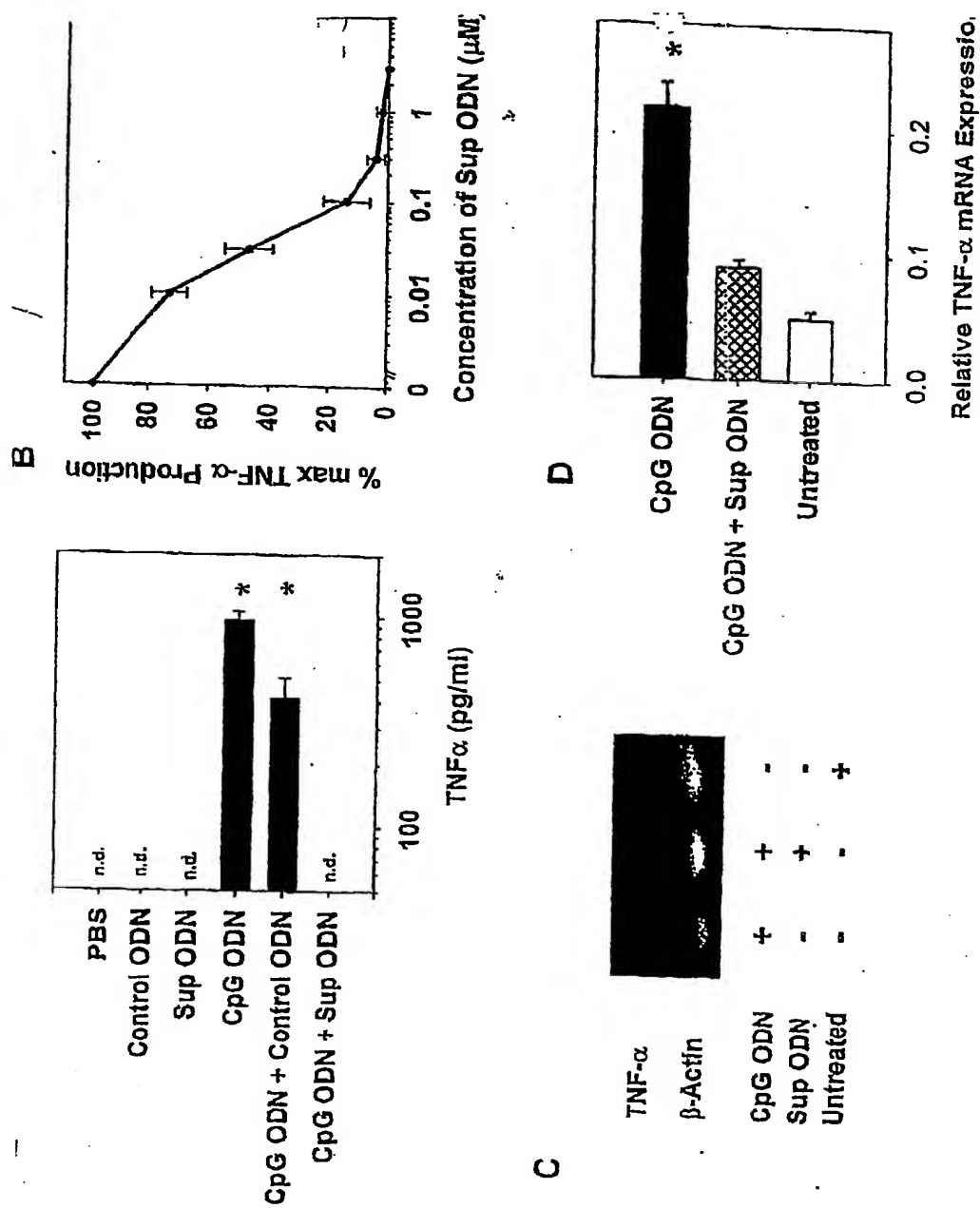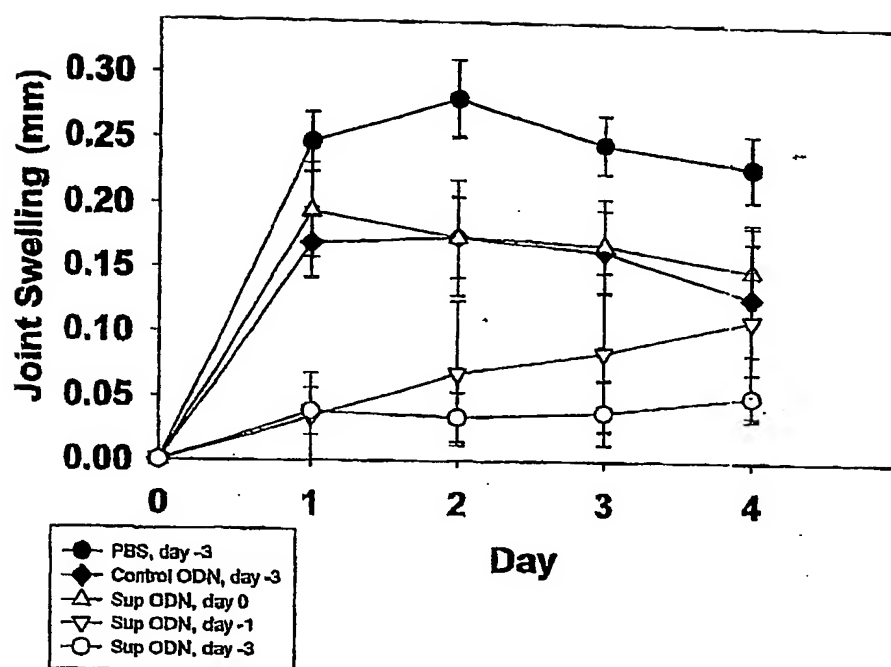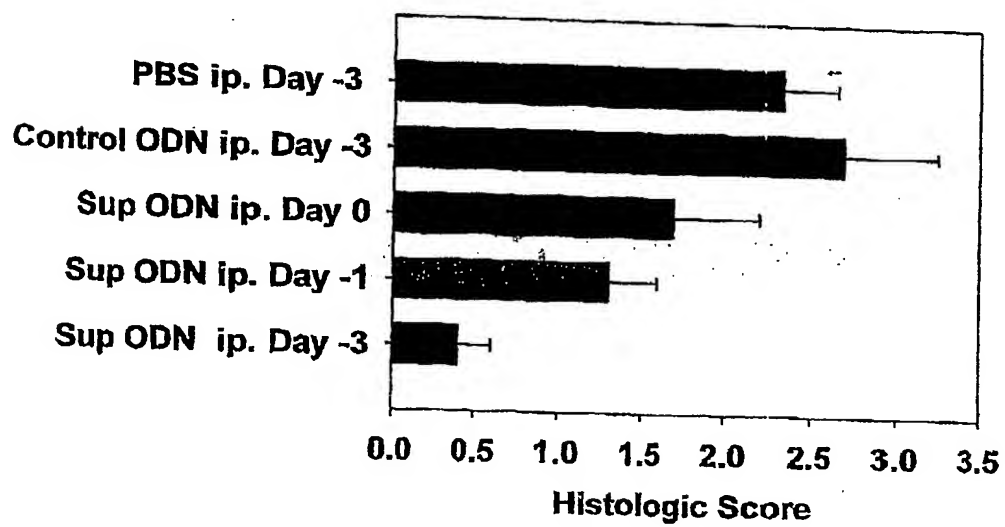
Fig. 13

Fig. 14

Systemic application of Sup ODN reduces CpG-Arthritis

Fig. 15

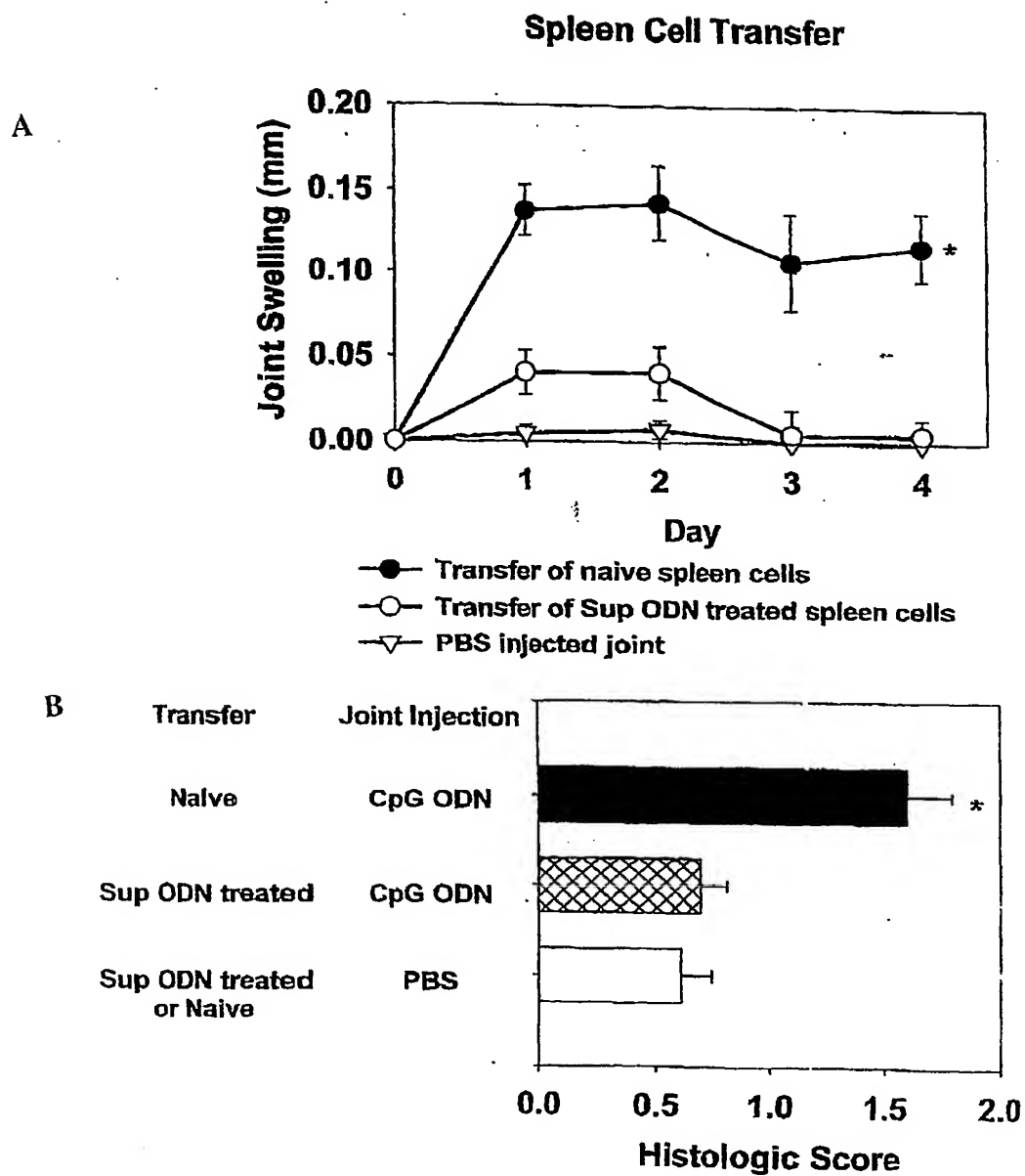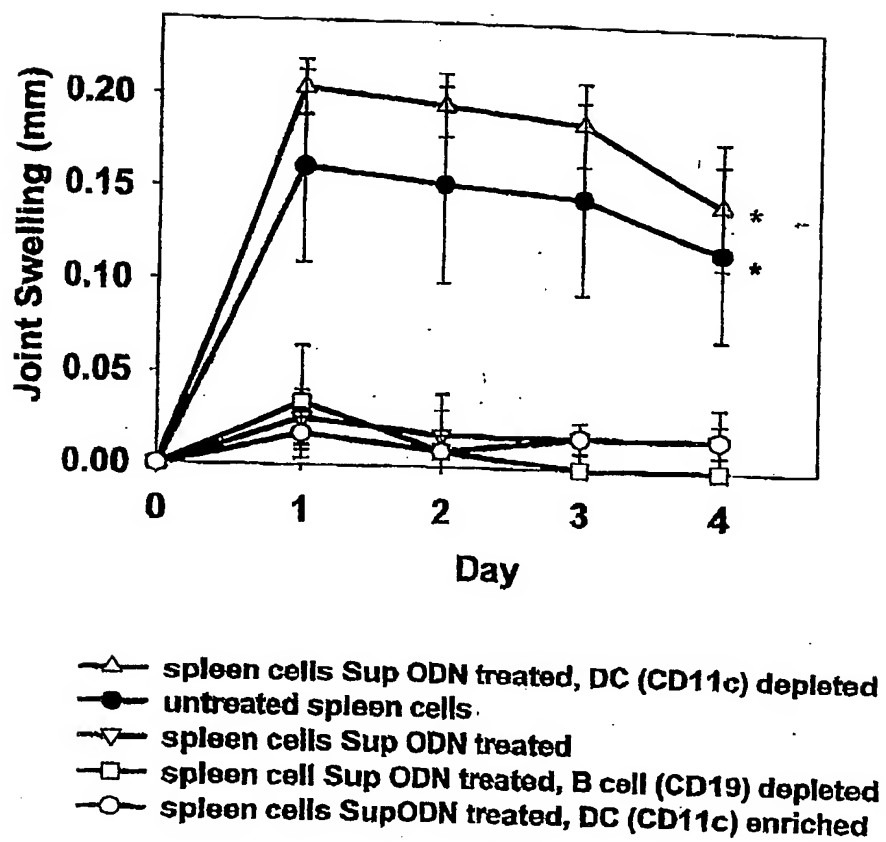Systemic effect of Sup ODN on CpG-induced Arthritis

A

Legend:
—●— Control ODN ip. day -3
—△— Sup ODN ip. day 0
—▽— Sup ODN ip. day -3

B

Fig. 16

Spleen Cell Transfer

A

- ● Transfer of naive spleen cells
- ○ Transfer of Sup ODN treated spleen cells
- ▽ PBS injected joint

B
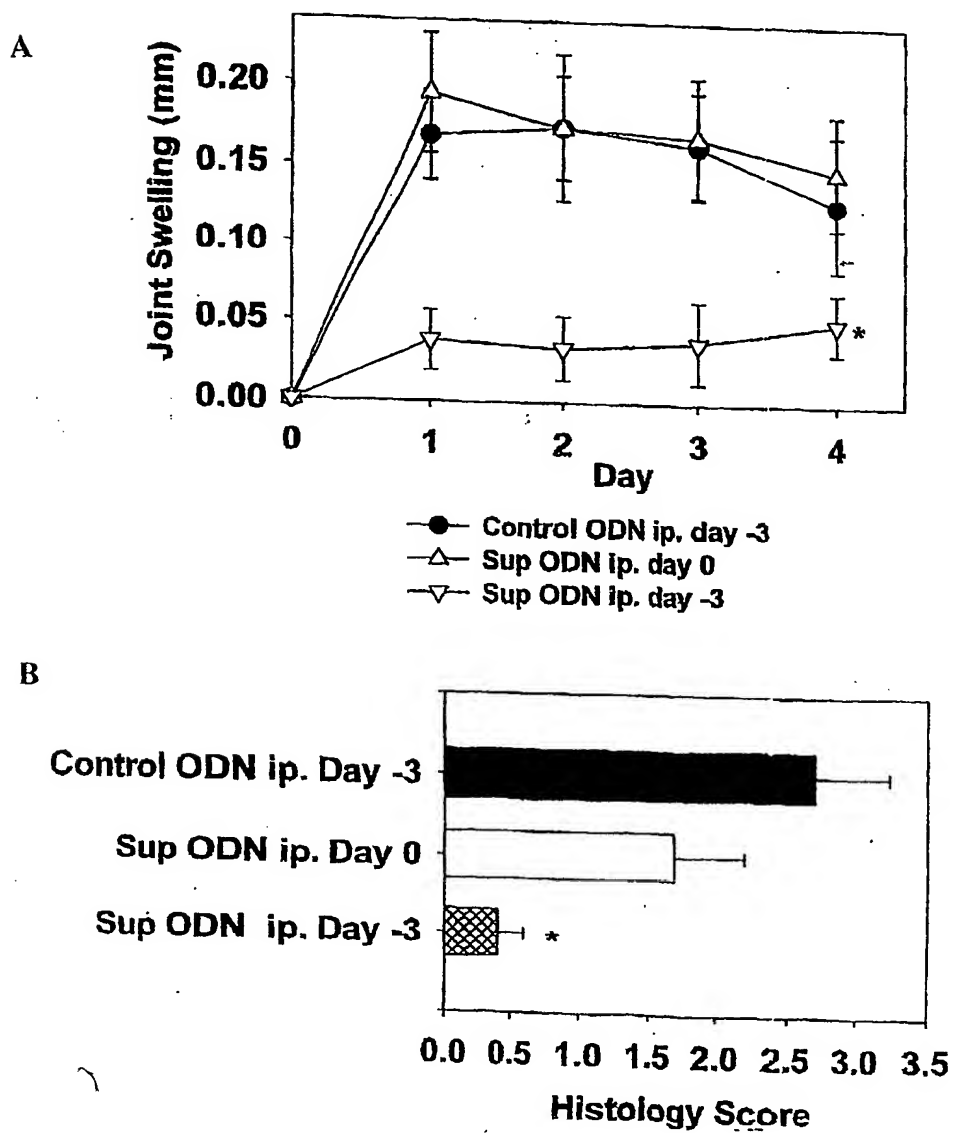
Fig. 17

Fig. 18
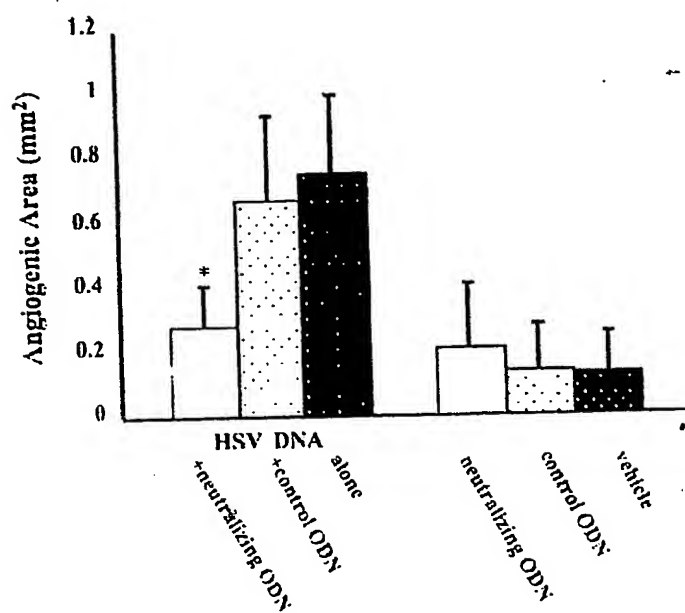
Fig. 19

SEQUENCE LISTING

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DEPARTMENT OF HEALTH AND HUMAN SERVICES
       Klinman, Dennis M.
       Gursel, Mayda M.
       Zeuner, Rainald
       Verthelyi, Daniela
       Gursel, Ihsan

<120> SUPPRESSORS OF CpG OLIGONUCLEOTIDES AND METHODS OF USE

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<150> US 60/324,484
<151> 2001-09-24

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<151> 2002-08-01

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STATEMENT ACCOMPANYING SEQUENCE LISTING

The sequence listing does not include matter that goes beyond the disclosure in the international application.

The printout of the attached Sequence Listing is identical to the computer readable sequence listing on the enclosed computer disk.

(54) Title: SUPPRESSORS OF CpG OLIGONUCLEOTIDES AND METHODS OF USE

(57) Abstract: The present disclosure relates to oligodeoxynucleotides that suppress an immune response. Methods are disclosed
for preventing or treating an immune-mediated disorder, such as, but not limited to, an autoimmune disease, by administering a ther-
apeutically effective amount of a suppressive oligodeoxynucleotide. Also disclosed are methods of suppressing an immune response
in a subject by administering a therapeutically effective amount of a suppressive oligodeoxynucleotide.

INTERNATIONAL SEARCH REPORT

A.	**CLASSIFICATION OF SUBJECT MATTER**

IPC(7) : C12Q 1/68; A01N 43/04; C07H 21/04; A61K 31/07
US CL : 435/6, 325, 375, 91.1; 536/23.1, 24.5, 24.3; 514/44

According to International Patent Classification (IPC) or to both national classification and IPC

B.	**FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)
  U.S. : 435/6, 325, 375, 91.1; 536/23.1, 24.5, 24.3; 514/44

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
Please See Continuation Sheet

C.	**DOCUMENTS CONSIDERED TO BE RELEVANT**

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	YAMADA et al. Effect of Suppressive DNA on CpG-Induced Immune Activation. Journal of Immunology, 2002, Vol. 169, pages 5590-5594, see entire article.	1-67
A	ZEUNER et al. Reduction of CpG-Induced Arthritis by Suppressive Oligonucleotides. Arthritis and Rheumatism, 2002, Vol. 46, No. 8 pages 2219-2224, see entire article.	1-67
A	STUNZ et al. Inhibitory oligonucleotides sepcifically block effects of stimulatory CpG oligonucletodies in B cells. European Journal of Immunology, 2002, Vol. 32, pages 1212-1222, see entire article.	1-67
X	KRIEG et al. Sequence motifs in adenoviral DNA block immune activation by stimulatory CpG motifs. Proc. Natl. Acad. Sci., 1998, Vol. 95, pp. 12631-12636, see especially Table 4.	1-67

☐ Further documents are listed in the continuation of Box C.   ☐ See patent family annex.

*	Special categories of cited documents:
"A"	document defining the general state of the art which is not considered to be of particular relevance
"E"	earlier application or patent published on or after the international filing date
"L"	document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
"O"	document referring to an oral disclosure, use, exhibition or other means
"P"	document published prior to the international filing date but later than the priority date claimed
"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"&"	document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
27 May 2003 (27.05.2003)	3 0 JUN 2003

Name and mailing address of the ISA/US	Authorized officer
Mail Stop PCT, Attn: ISA/US Commissioner for Patents P.O. Box 1450 Alexandria, Virginia 22313-1450	Terra C. Gibbs Felicia D. Roberts for
Facsimile No. (703)305-3230	Telephone No. (703) 308-0196

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Continuation of B. FIELDS SEARCHED Item 3:
CaPlus, Embase, Medline, CancerLit,
search terms: Inhibitory oligonucleotides, CpG oligonucleotides, suppressive oligonucleotides